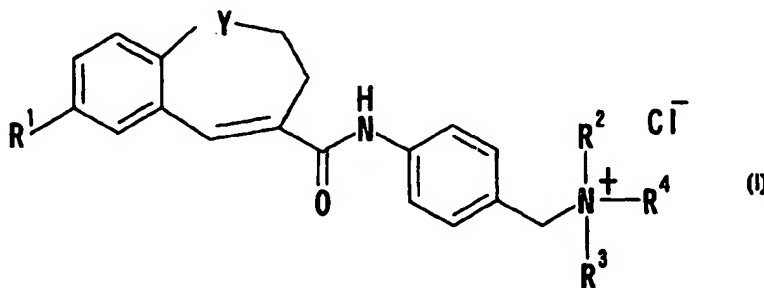




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(54) Title: QUATERNARY AMMONIUM SALTS AND THEIR USE



(57) Abstract

This invention is to provide a compound for antagonizing CCR5, said compound being represented by formula (I), wherein R¹ is an optionally substituted phenyl or an optionally substituted thienyl; Y is -CH₂-, -S- or -O-; and R², R³ and R⁴ are independently an optionally substituted aliphatic hydrocarbon group or an optionally substituted alicyclic heterocyclic ring group, and being effective for the prevention and treatment of infectious disease of HIV.

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DESCRIPTION

Quaternary Ammonium Salts and Their Use

5 Technical Field

The present invention relates to a quaternary ammonium salt which has CCR5 antagonistic activity and which is used for the treatment or prevention of infectious disease of HIV, etc.

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Background Art

Recently, HIV (human immunodeficiency virus) protease inhibitors are developed for method of the treatment of AIDS (acquired immunological deficient syndrome) and use of the protease inhibitors in combination with conventional two HIV reverse transcriptase inhibitors provides with a further progress of the treatment of AIDS. However, these drugs and their combination use are not sufficient for the eradication of AIDS, and development of new anti-AIDS drugs having different activity and mechanism are sought for.

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As a receptor from which HIV invades to a target cell, CD4 is so far known, and recently CCR5 as a second receptor of macrophage-tropic HIV and CXCR4 as a second receptor of T cell-tropic HIV, each of which is G protein-coupled chemokine receptor having seven transmembrane domains, are respectively found out. These chemokine receptors are thought to play an essential role in establishment and spread of HIV infection. In fact, it is reported that a person who is resistant to HIV infection in spite of several exposures retains mutation of homo deletion of CCR5 gene. Therefore, a CCR5 antagonist is expected to be a new anti-HIV drug. However, so far, there has been no report that a CCR5 antagonist is developed as a therapeutic agent of AIDS.

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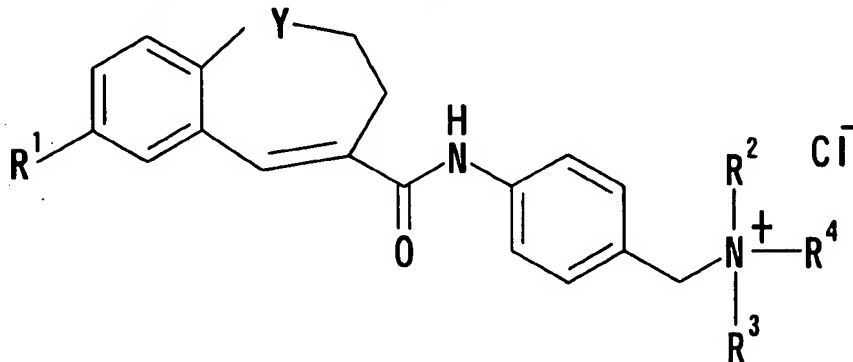
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Disclosure of Invention

The present invention is to provide a quaternary ammonium salt having CCR5 antagonistic activity and less toxicity; and a composition for antagonizing CCR5 (a drug for the treatment or prevention of infectious disease of HIV (in particular, AIDS), etc.) comprising said quaternary ammonium salt.

The present inventors diligently made extensive studies on compounds having CCR5 antagonistic activity and less toxicity, as a result, they found that a quaternary ammonium salt of the following formula (I) [hereinafter, referred to as Compound (I)] unexpectedly possesses potent CCR5 antagonistic activity and clinically desirable pharmaceutical effect (e.g. remarkable inhibition of HIV infection to human peripheral mononuclear cells, etc.) and also that Compound (I) has superior solubility, physicochemical properties (stability, anti-coloring effect, etc.), etc. and is useful for applying as an injection. Based on the finding, the present invention was accomplished.

More specifically, the present invention relates to (1) a compound of the formula:



wherein R¹ is an optionally substituted phenyl or an optionally substituted thienyl; Y is -CH₂-, -S- or -O-; and R², R³ and R⁴ are independently an optionally substituted aliphatic hydrocarbon group or an optionally substituted alicyclic heterocyclic ring group; or a pro-drug thereof;

- (2) a compound of the above (1), wherein R^2 and R^3 are independently an optionally substituted acyclic hydrocarbon group;
- (3) a compound of the above (1), wherein R^2 and R^3 are independently an optionally substituted alkyl group;
- (4) a compound of the above (1), wherein R^4 is an optionally substituted alicyclic hydrocarbon group or an optionally substituted alicyclic heterocyclic ring group;
- (5) a compound of the above (4), wherein the alicyclic hydrocarbon group is cycloalkyl;
- (6) a compound of the above (4), wherein the alicyclic hydrocarbon group is cyclohexyl;
- (7) a compound of the above (4), wherein the alicyclic heterocyclic ring group is a saturated alicyclic heterocyclic ring group;
- (8) a compound of the above (4), wherein the alicyclic heterocyclic ring group is tetrahydropyranyl, tetrahydrothiopyranyl or piperidyl;
- (9) N,N-Dimethyl-N-(4-(((2-(4-methylphenyl)-6,7-dihydro-5H-benzocyclohepten-8-yl)carbonyl)amino)benzyl)-N-(4-tetrahydropyranyl)ammonium chloride;
- (10) N,N-Dimethyl-N-(((7-(4-methylphenyl)-2,3-dihydro-1-benzoxepin-4-yl)carbonyl)amino)benzyl)-N-(4-oxocyclohexyl)ammonium chloride;
- (11) N-(4-(((7-(4-Ethoxyphenyl)-2,3-dihydro-1-benzoxepin-4-yl)carbonyl)amino)benzyl)-N,N-dimethyl-N-(4-tetrahydropyranyl)ammonium chloride;
- (12) a pharmaceutical composition which comprises a compound of the above (1);
- (13) a pharmaceutical composition for antagonizing CCR5 which comprises a compound of the above (9);
- (14) a composition of the above (13), which is for the treatment or prevention of infectious disease of HIV;
- (15) a composition of the above (13), which is for the treatment or prevention of AIDS;

- (16) a composition of the above (13), which is for the prevention of the progression of AIDS;
- (17) a composition of the above (13), which is used in combination with a protease inhibitor and/or a reverse transcriptase inhibitor;
- (18) a composition of the above (17), wherein the reverse transcriptase inhibitor is zidovudine, didanosine, zalcitabine, lamivudine, stavudine, nevirapine, delavirdine, efavirenz or abacavir;
- (19) a composition of the above (17), wherein the protease inhibitor is saquinavir, ritonavir, indinavir or nelfinavir;
- (20) use of the compound of the above (9) in combination with a protease inhibitor and/or a reverse transcriptase inhibitor for the treatment or prevention of infectious disease of HIV; etc.

In the above formula (I), example of the "substituents" which the "phenyl group" and the "thienyl group" of the "optionally substituted phenyl group" and "optionally substituted thienyl group" represented by R^1 may have include halogen atom, nitro, cyano, an optionally substituted alkyl, an optionally substituted cycloalkyl, an optionally substituted hydroxy group, an optionally substituted thiol group, an optionally substituted amino group, an optionally substituted acyl, an optionally esterified carboxyl group, an optionally substituted aromatic group, etc.

Examples of the halogen as the substituents for R^1 include fluorine, chlorine, bromine, iodine, etc. Among others, fluorine and chlorine are preferable.

Examples of the alkyl in the optionally substituted alkyl as the substituents for R^1 include a straight or branched C_{1-10} alkyl such as methyl, ethyl, propyl, isopropyl, butyl, isobutyl, sec-butyl, tert-butyl, pentyl, isopentyl, neopentyl, hexyl, heptyl, octyl, nonyl, decyl, etc., and preferably lower (C_{1-6}) alkyl.

Examples of the substituents in the optionally

substituted alkyl include halogen (e.g. fluorine, chlorine, bromine, iodine, etc.), nitro, cyano, hydroxy group, thiol group, amino group, carboxyl group, an optionally halogenated C₁₋₄ alkoxy (e.g. methoxy, ethoxy, trifluoromethoxy, trifluoroethoxy, etc.), C₁₋₄ alkanoyl (e.g. acetyl, propionyl, etc.), C₁₋₄ alkylsulfonyl (e.g. methanesulfonyl, ethanesulfonyl, etc.), etc., and the number of the substituents are preferably 1 to 3.

Examples of the cycloalkyl in the optionally substituted cycloalkyl as the substituents for R¹ include C₃₋₇ cycloalkyl, etc. such as cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl, etc.

Examples of the substituents in the optionally substituted cycloalkyl include halogen (e.g. fluorine, chlorine, bromine, iodine, etc.), nitro, cyano, hydroxy group, thiol group, amino group, carboxyl group, an optionally halogenated C₁₋₄ alkyl (e.g. trifluoromethyl, methyl, ethyl, etc.), an optionally halogenated C₁₋₄ alkoxy (e.g. methoxy, ethoxy, trifluoromethoxy, trifluoroethoxy, etc.), C₂₋₄ alkanoyl (e.g. acetyl, propionyl, etc.), C₁₋₄ alkylsulfonyl (e.g. methanesulfonyl, ethanesulfonyl, etc.), etc., and the number of the substituents are preferably 1 to 3.

Examples of the substituents in the optionally substituted hydroxy group as the substituents for R¹ include (1) an optionally substituted alkyl (e.g. C₁₋₁₀ alkyl such as methyl, ethyl, propyl, isopropyl, butyl, isobutyl, sec-butyl, tert-butyl, pentyl, isopentyl, neopentyl, hexyl, heptyl, octyl, nonyl, decyl, etc., preferably lower (C₁₋₆) alkyl, etc.); (2) an optionally substituted cycloalkyl (e.g. C₃₋₇ cycloalkyl, etc. such as cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl, etc.); (3) an optionally substituted alkenyl (e.g. C₂₋₁₀ alkenyl such as allyl, crotyl, 2-pentenyl, 3-hexenyl, etc., preferably lower (C₁₋₆) alkenyl, etc.);

- (4) an optionally substituted cycloalkenyl (e.g. C₃₋₇ cycloalkenyl, etc. such as 2-cyclopentenyl, 2-cyclohexenyl, 2-cyclopentenylmethyl, 2-cyclohexenylmethyl, etc.);
- (5) an optionally substituted aralkyl (e.g. phenyl-C₁₋₄ alkyl (e.g. benzyl, phenethyl, etc.), etc.);
- (6) an optionally substituted acyl (e.g. C₂₋₄ alkanoyl (e.g. acetyl, propionyl, butyryl, isobutyryl, etc.), C₁₋₄ alkylsulfonyl (e.g. methanesulfonyl, ethanesulfonyl, etc.), etc.);
- (7) an optionally substituted aryl (e.g. phenyl, naphthyl, etc.); etc.

Examples of the substituents which the above-mentioned (1) optionally substituted alkyl, (2) optionally substituted cycloalkyl, (3) optionally substituted alkenyl, (4) optionally substituted cycloalkenyl, (5) optionally substituted aralkyl, (6) optionally substituted acyl and (7) optionally substituted aryl may have include halogen (e.g. fluorine, chlorine, bromine, iodine, etc.), nitro, cyano, hydroxy group, thiol group, amino group, carboxyl group, an optionally halogenated C₁₋₄ alkyl (e.g. trifluoromethyl, methyl, ethyl, etc.), an optionally halogenated C₁₋₄ alkoxy (e.g. methoxy, ethoxy, trifluoromethoxy, trifluoroethoxy, etc.), C₂₋₄ alkanoyl (e.g. acetyl, propionyl, etc.), C₁₋₄ alkylsulfonyl (e.g. methanesulfonyl, ethanesulfonyl, etc.), etc., and the number of the substituents are preferably 1 to 3.

Examples of the substituents in the optionally substituted thiol group as the substituents for R¹ are the same as the above-described substituents in the optionally substituted hydroxy group as the substituents for R¹, and among others,

- (1) an optionally substituted alkyl (e.g. C₁₋₁₀ alkyl such as methyl, ethyl, propyl, isopropyl, butyl, isobutyl, sec-butyl, tert-butyl, pentyl, isopentyl, neopentyl, hexyl, heptyl, octyl, nonyl, decyl, etc., preferably lower (C₁₋₆) alkyl, etc.);

(2) an optionally substituted cycloalkyl (e.g. C₃₋₇ cycloalkyl, etc. such as cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl, etc.);

(3) an optionally substituted aralkyl (e.g. phenyl-C₁₋₄ alkyl (e.g. benzyl, phenethyl, etc.), etc.);

(4) an optionally substituted aryl (e.g. phenyl, naphthyl, etc.); etc. are preferable.

Examples of the substituents which the above-mentioned (1) optionally substituted alkyl, (2) optionally substituted cycloalkyl, (3) optionally substituted aralkyl and (4) optionally substituted aryl may have include halogen (e.g. fluorine, chlorine, bromine, iodine, etc.), nitro, cyano, hydroxy group, thiol group, amino group, carboxyl group, an optionally halogenated C₁₋₄ alkyl (e.g. trifluoromethyl, methyl, ethyl, etc.), an optionally halogenated C₁₋₄ alkoxy (e.g. methoxy, ethoxy, trifluoromethoxy, trifluoroethoxy, etc.), C₂₋₄ alkanoyl (e.g. acetyl, propionyl, etc.), C₁₋₄ alkylsulfonyl (e.g. methanesulfonyl, ethanesulfonyl, etc.), etc., and the number of the substituents are preferably 1 to 3.

Examples of the substituents in the optionally substituted amino group as the substituents for R¹ are the same as the above-described substituents in the optionally substituted hydroxy group as the substituents for R¹, and examples of the optionally substituted amino group as the substituents for R¹ include an amino group which may have one to two substituents selected from the above-described substituents in the optionally substituted hydroxy group as the substituents for R¹, etc. Among others, as the substituents in the optionally substituted amino group as the substituents for R¹,

(1) an optionally substituted alkyl (e.g. C₁₋₁₀ alkyl such as methyl, ethyl, propyl, isopropyl, butyl, isobutyl, sec-butyl, tert-butyl, pentyl, isopentyl, neopentyl, hexyl, heptyl, octyl, nonyl, decyl, etc., preferably lower (C₁₋₆) alkyl, etc.);

- (2) an optionally substituted cycloalkyl (e.g. C₃, cycloalkyl, etc. such as cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl, etc.);
- (3) an optionally substituted alkenyl (e.g. C₂₋₁₀ alkenyl such as allyl, crotyl, 2-pentenyl, 3-hexenyl, etc., preferably lower (C₁₋₆) alkenyl, etc.);
- (4) an optionally substituted cycloalkenyl (e.g. C₃, cycloalkenyl, etc. such as 2-cyclopentenyl, 2-cyclohexenyl, 2-cyclopentenylmethyl, 2-cyclohexenylmethyl, etc.);
- (5) an optionally substituted acyl (e.g. C₂₋₄ alkanoyl (e.g. acetyl, propionyl, butyryl, isobutyryl, etc.), C₁₋₄ alkylsulfonyl (e.g. methanesulfonyl, ethanesulfonyl, etc.), etc.);
- (6) an optionally substituted aryl (e.g. phenyl, naphthyl, etc.); etc. are preferable.

Examples of the substituents, which each of the above-described (1) optionally substituted alkyl, (2) optionally substituted cycloalkyl, (3) optionally substituted alkenyl, (4) optionally substituted cycloalkenyl, (5) optionally substituted acyl and (6) optionally substituted aryl may have, include halogen (e.g. fluorine, chlorine, bromine, iodine, etc.), nitro, cyano, hydroxy group, thiol group, amino group, carboxyl group, an optionally halogenated C₁₋₄ alkyl (e.g. trifluoromethyl, methyl, ethyl, etc.), an optionally halogenated C₁₋₄ alkoxy (e.g. methoxy, ethoxy, trifluoromethoxy, trifluoroethoxy, etc.), C₂₋₄ alkanoyl (e.g. acetyl, propionyl, etc.), C₁₋₄ alkylsulfonyl (e.g. methanesulfonyl, ethanesulfonyl, etc.), etc., and the number of the substituents are preferably 1 to 3.

The substituents in the optionally substituted amino group as the substituents for R¹ may bind to each other to form a cyclic amino group (e.g. 5- to 6-membered cyclic amino, etc. such as pyrrolidine, pyrroline, piperazine, piperidine, morpholine, thiomorpholine, pyrrole, imidazole, etc.). Said cyclic amino group may have a substituent, and examples

of the substituents include halogen (e.g. fluorine, chlorine, bromine, iodine, etc.), nitro, cyano, hydroxy group, thiol group, amino group, carboxyl group, an optionally halogenated C₁₋₄ alkyl (e.g. trifluoromethyl, methyl, ethyl, etc.), an optionally halogenated C₁₋₄ alkoxy (e.g. methoxy, ethoxy, trifluoromethoxy, trifluoroethoxy, etc.), C₂₋₄ alkanoyl (e.g. acetyl, propionyl, etc.), C₁₋₄ alkylsulfonyl (e.g. methanesulfonyl, ethanesulfonyl, etc.), etc., and the number of the substituents are preferably 1 to 3.

10 Examples of the optionally substituted acyl as the substituents for R¹ include a carbonyl group or a sulfonyl group binding to

(1) hydrogen;

15 (2) an optionally substituted alkyl (e.g. C₁₋₁₀ alkyl such as methyl, ethyl, propyl, isopropyl, butyl, isobutyl, sec-butyl, tert-butyl, pentyl, isopentyl, neopentyl, hexyl, heptyl, octyl, nonyl, decyl, etc., preferably lower (C₁₋₆) alkyl, etc.);

20 (3) an optionally substituted cycloalkyl (e.g. C₃₋₇ cycloalkyl, etc. such as cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl, etc.);

(4) an optionally substituted alkenyl (e.g. C₂₋₁₀ alkenyl such as allyl, crotyl, 2-pentenyl, 3-hexenyl, etc., preferably lower (C₂₋₆) alkenyl, etc.);

25 (5) an optionally substituted cycloalkenyl (e.g. C₃₋₇ cycloalkenyl, etc. such as 2-cyclopentenyl, 2-cyclohexenyl, 2-cyclopentenylmethyl, 2-cyclohexenylmethyl, etc.);

(6) an optionally substituted 5- to 6-membered monocyclic aromatic group (e.g. phenyl, pyridyl, etc.); etc.

30 Examples of the acyl include acetyl, propionyl, butyryl, isobutyryl, valeryl, isovaleryl, pivaloyl, hexanoyl, heptanoyl, octanoyl, cyclobutanecarbonyl, cyclopentanecarbonyl, cyclohexanecarbonyl, cycloheptanecarbonyl, crotonyl, 2-cyclohexenecarbonyl, benzoyl, nicotinoyl, methanesulfonyl, ethanesulfonyl, etc.

35 Examples of the substituents, which the above-

mentioned (2) optionally substituted alkyl, (3) optionally substituted cycloalkyl, (4) optionally substituted alk nyl, (5) optionally substituted cycloalkenyl and (6) optionally substituted 5- to 6-membered monocyclic aromatic group may
5 have, include halogen (e.g. fluorine, chlorine, bromine, iodine, etc.), nitro, cyano, hydroxy group, thiol group, amino group, carboxyl group, an optionally halogenated C₁₋₄ alkyl (e.g. trifluoromethyl, methyl, ethyl, etc.), an optionally halogenated C₁₋₄ alkoxy (e.g. methoxy, ethoxy,
10 trifluoromethoxy, trifluoroethoxy, etc.), C₂₋₄ alkanoyl (e.g. acetyl, propionyl, etc.), C₁₋₄ alkylsulfonyl (e.g. methanesulfonyl, ethanesulfonyl, etc.), etc., and the number of the substituents are preferably 1 to 3.

Examples of the optionally esterified carboxyl group
15 as the substituents for R¹ include a carbonyloxy group binding to

- (1) hydrogen;
- (2) an optionally substituted alkyl (e.g. C₁₋₁₀ alkyl such as methyl, ethyl, propyl, isopropyl, butyl, isobutyl,
20 sec-butyl, tert-butyl, pentyl, isopentyl, neopentyl, hexyl, heptyl, octyl, nonyl, decyl, etc., preferably lower (C₁₋₆) alkyl, etc.);
- (3) an optionally substituted cycloalkyl (e.g. C₃₋₇ cycloalkyl, etc. such as cyclopropyl, cyclobutyl,
25 cyclopentyl, cyclohexyl, cycloheptyl, etc.);
- (4) an optionally substituted alkenyl (e.g. C₂₋₁₀ alkenyl such as allyl, crotyl, 2-pentenyl, 3-hexenyl, etc., preferably lower (C₂₋₆) alkenyl, etc.);
- (5) an optionally substituted cycloalkenyl (e.g. C₃₋₇ cycloalkenyl, etc. such as 2-cyclopentenyl, 2-cyclohexenyl, 2-cyclopentenylmethyl, 2-cyclohexenylmethyl, etc.);
- (6) an optionally substituted aryl (e.g. phenyl, naphthyl, etc.); etc., and preferably carboxyl, lower (C₁₋₆) alkoxy-carbonyl, aryloxy-carbonyl (e.g. methoxycarbonyl,
30 ethoxycarbonyl, propoxycarbonyl, phenoxycarbonyl, naphthoxycarbonyl, etc.), etc.

Examples of the substituents, which the above-mentioned (2) optionally substituted alkyl, (3) optionally substituted cycloalkyl, (4) optionally substituted alkenyl, (5) optionally substituted cycloalkenyl and (6) optionally substituted aryl may have, include halogen (e.g. fluorine, chlorine, bromine, iodine, etc.), nitro, cyano, hydroxy group, thiol group, amino group, carboxyl group, an optionally halogenated C₁₋₄ alkyl (e.g. trifluoromethyl, methyl, ethyl, etc.), an optionally halogenated C₁₋₄ alkoxy (e.g. methoxy, ethoxy, trifluoromethoxy, trifluoroethoxy, etc.), C₂₋₄ alkanoyl (e.g. acetyl, propionyl, etc.), C₁₋₄ alkylsulfonyl (e.g. methanesulfonyl, ethanesulfonyl, etc.), etc., and the number of the substituents are preferably 1 to 3.

Examples of the aromatic group in the optionally substituted aromatic group as the substituents for R¹ include 5- to 6-membered homocyclic or heterocyclic ring aromatic ring, etc. such as phenyl, pyridyl, furyl, thienyl, pyrrolyl, imidazolyl, pyrazolyl, thiazolyl, oxazolyl, isothiazolyl, isoxazolyl, tetrazolyl, pyrazinyl, pyrimidinyl, pyridazinyl, triazolyl, etc.

Examples of the substituents for these aromatic group include halogen (e.g. fluorine, chlorine, bromine, iodine, etc.), nitro, cyano, hydroxy group, thiol group, amino group, carboxyl group, an optionally halogenated C₁₋₄ alkyl (e.g. trifluoromethyl, methyl, ethyl, etc.), an optionally halogenated C₁₋₄ alkoxy (e.g. methoxy, ethoxy, trifluoromethoxy, trifluoroethoxy, etc.), C₂₋₄ alkanoyl (e.g. acetyl, propionyl, etc.), C₁₋₄ alkylsulfonyl (e.g. methanesulfonyl, ethanesulfonyl, etc.), etc., and the number of the substituents are preferably 1 to 3.

The number of the above-mentioned substituents for R¹ is 1-4 (preferably 1-2) and they may be same or different and present at any possible position on the ring represented by R¹. When two or more substituents are present on the "phenyl group" and the "thienyl group" of the "optionally

substituted phenyl group" and "optionally substituted thienyl group" represented by R^1 , two substituents among them may bind to each other to form a lower (C_{1-6}) alkylene (e.g. trimethylene, tetramethylene, etc.), a lower (C_{1-6}) alkyleneoxy (e.g. $-CH_2-O-CH_2-$, $-O-CH_2-CH_2-$, etc.), a lower (C_{1-6}) alkylenedioxy (e.g. $-O-CH_2-O-$, $-O-CH_2-CH_2-O-$, etc.), a lower (C_{1-6}) alkenylene (e.g. $-CH_2-CH=CH-$, $-CH_2-CH_2-CH=CH-$, $-CH_2-CH=CH-CH_2-$, etc.), a lower (C_{1-6}) alkadienylene (e.g. $-CH=CH-CH=CH-$, etc.), etc.

Preferred examples of the "substituents", which the "phenyl group" and the "thienyl group" of the "optionally substituted phenyl group" and "optionally substituted thienyl group" represented by R^1 may have, include an optionally halogenated lower (C_{1-4}) alkyl (e.g. methyl, ethyl, t-butyl, trifluoromethyl, etc.), an optionally halogenated lower (C_{1-4}) alkoxy (e.g. methoxy, ethoxy, t-butoxy, trifluoromethoxy, etc.), halogen (e.g. fluorine, chlorine, etc.), nitro, cyano, an amino group optionally substituted with 1-2 lower (C_{1-4}) alkyl groups (e.g. amino, methylamino, dimethylamino, etc.), 5- to 6-membered cyclic amino (e.g. 1-pyrrolidinyl, 1-piperazinyl, 1-piperidinyl, 4-morpholino, 4-thiomorpholino, 1-imidazolyl, 4-tetrahydropyranyl, etc.), etc., and when R^1 is a phenyl group, the "substituent" is preferably present at para position.

In the above formula (I), Y is $-CH_2-$, $-S-$ or $-O-$; and preferably $-CH_2-$ or $-O-$.

In the above formula (I), R^2 , R^3 and R^4 are independently an optionally substituted aliphatic hydrocarbon group or an optionally substituted alicyclic heterocyclic ring group.

Examples of the "aliphatic hydrocarbon group" in the "optionally substituted aliphatic hydrocarbon group" represented by R^2 , R^3 and R^4 include (1) an optionally substituted alkyl (e.g. C_{1-10} alkyl such as methyl, ethyl, propyl, isopropyl, butyl, isobutyl, sec-butyl, tert-butyl, pentyl, isopentyl, neopentyl, hexyl,

heptyl, octyl, nonyl, decyl, etc., preferably lower (C_{1-6}) alkyl, etc.);

(2) an optionally substituted cycloalkyl (e.g. C_{3-8} cycloalkyl, etc. such as cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl, cyclooctyl, etc.); provided that

(3) an optionally substituted alkenyl (e.g. C_{2-10} alkenyl such as allyl, crotyl, 2-pentenyl, 3-hexenyl, etc., preferably lower (C_{2-6}) alkenyl, etc.);

(4) an optionally substituted cycloalkenyl (e.g. C_{3-7} cycloalkenyl, etc. such as 2-cyclopentenyl, 2-cyclohexenyl, 2-cyclopentenylmethyl, 2-cyclohexenylmethyl, etc.); etc.

Examples of the "alicyclic heterocyclic ring group" in the "optionally substituted alicyclic heterocyclic ring group" represented by R^2 , R^3 and R^4 include a 5- to 6-membered non-aromatic heterocyclic ring containing 1 to 4 hetero-atoms consisting of 1 to 2 kinds of hetero-atoms selected from oxygen atom, sulfur atom and nitrogen atom such as tetrahydrofuran, tetrahydrothiophene, dithiolane, oxathiolane, pyrrolidine, pyrroline, imidazolidine, imidazoline, pyrazolidine, pyrazoline, piperidine, piperazine, oxazine, oxadiazine, thiazine, thiadiazine, morpholine, thiomorpholine, pyran, tetrahydropyran, etc.; preferably a saturated 5- to 6-membered heterocyclic ring, etc., such as tetrahydrofuran, piperidine, tetrahydropyran, tetrahydrothiopyran, etc.); etc.

Examples of the substituents, which the "aliphatic hydrocarbon group" and the "alicyclic heterocyclic ring group" in the "optionally substituted aliphatic hydrocarbon group" and the "optionally substituted alicyclic heterocyclic ring group" represented by R^2 , R^3 and R^4 may have, include halogen (e.g. fluorine, chlorine, bromine, iodine, etc.), an optionally halogenated lower (C_{1-4}) alkyl, an optionally halogenated C_{1-4} alkoxy (e.g. methoxy, ethoxy, trifluoromethoxy, trifluoroethoxy, etc.), C_{1-4} alkanoyl (e.g. acetyl, propionyl, etc.), C_{1-4} alkylsulfonyl (e.g.

methanesulfonyl, ethanesulfonyl, etc.), phenyl, phenyl-lower (C₁₋₄) alkyl, C₃, cycloalkyl, cyano, nitro, oxo, hydroxy group, thiol group, amino group, carboxyl group, lower (C₁₋₄) alkoxy-carbonyl (preferably, halogen, an optionally
5 halogenated lower (C₁₋₄) alkyl, an optionally halogenated lower (C₁₋₄) alkoxy, phenyl-lower (C₁₋₄) alkyl, C₃, cycloalkyl, cyano, oxo, hydroxy group, etc.), etc., and the number of the substituents are preferably 1 to 3.

In the above formula (I), as the group R² or R³, an
10 optionally substituted acyclic hydrocarbon group is preferable, an optionally substituted alkyl group is more preferable. In particular, the groups R² and R³ are preferably the same and more preferably both of the groups R² and R³ are methyl.

15 In the above formula (I), as the group R⁴, an optionally substituted alicyclic hydrocarbon group or an optionally substituted alicyclic heterocyclic ring group and more preferably an optionally substituted cycloalkyl group or an optionally substituted saturated alicyclic heterocyclic
20 ring group. In particular, R⁴ is preferably an optionally substituted cyclohexyl or an optionally substituted 6-membered saturated alicyclic heterocyclic ring group and more preferably an optionally substituted cycloalkyl, an optionally substituted tetrahydropyranyl, an optionally
25 substituted tetrahydrothiopyranyl or an optionally substituted piperidyl.

The compound of the formula (I) of the present invention may be hydrated or solvated. When the compound of the formula
30 (I) of the present invention exists as configuration isomer, diastereomer, conformer, etc., it is possible to isolate individual isomers with per se known separation and purification method, if desired. When the compound of the formula (I) of the present invention is racemate, it can be
35 separated into (S)-compound and (R)-compound with usual optical resolution and individual optical isomers and a

mixture thereof are included in the scope of the present invention.

The pro-drug of Compound (I) means a compound which is converted to Compound (I) under the physiological condition or with a reaction due to an enzyme, an gastric acid, etc. in the living body, that is, a compound which is converted to Compound (I) with oxidation, reduction, hydrolysis, etc. according to an enzyme; a compound which is converted to Compound (I) with gastric acid, etc.; etc.

Examples of the pro-drug of Compound (I) include a compound wherein an amino group of Compound (I) is substituted with acyl, alkyl, phosphoric acid, etc. (e.g. a compound wherein an amino group of Compound (I) is substituted with eicosanoyl, alanyl, pentylaminocarbonyl, (5-methyl-2-oxo-1,3-dioxolen-4-yl)methoxycarbonyl, tetrahydrofuranyl, pyrrolidylmethyl, pivaloyloxymethyl, tert-butyl, etc.); a compound wherein an hydroxy group of Compound (I) is substituted with acyl, alkyl, phosphoric acid, boric acid, etc. (e.g. a compound wherein an hydroxy group of Compound (I) is substituted with acetyl, palmitoyl, propanoyl, pivaloyl, succinyl, fumaryl, alanyl, dimethylaminomethylcarbonyl, etc.); a compound wherein a carboxyl group of Compound (I) is modified with ester, amide, etc. (e.g. a compound wherein a carboxyl group of Compound (I) is modified with ethyl ester, phenyl ester, carboxymethyl ester, dimethylaminomethyl ester, pivaloyloxymethyl ester, ethoxycarbonyloxyethyl ester, phthalidyl ester, (5-methyl-2-oxo-1,3-dioxolen-4-yl)methyl ester, cyclohexyloxycarbonylethyl ester, methyl amide, etc.); etc. These pro-drug can be produced by per se known method from Compound (I).

The pro-drug of Compound (I) may be a compound which is converted into Compound (I) under the physiological conditions as described in "Pharmaceutical Research and Development", Vol. 7 (Drug Design), pages 163-198 published in 1990 by Hirokawa Publishing Co. (Tokyo, Japan).

Compound (I) may be labeled with isotope (e.g. ^3H , ^{14}C , ^{35}S , ^{125}I , etc.) , etc.

The present compound of the formula (I) alone or as an admixture with a pharmaceutically acceptable carrier may be non-orally administered as a liquid preparation such as an injection.

Examples of the carriers include various organic or inorganic carriers which are generally used in this field. For example, a solvent, a solubilizer, a suspending agent, a isotonizing agent, a buffer, a soothing agent, etc. are used in the liquid formulations. In addition, if desired, an appropriate additive such as a preservative, an antioxidant, a colorant, a sweetener, etc. may be used in the above formulations.

Examples of the solvent include water for injection, alcohol, propyleneglycol, macrogol, sesame oil, corn oil, etc. Examples of the solubilizer include polyethyleneglycol, propyleneglycol, polyoxyethylene (20) sorbitan mono-oleate (polysorbate 80), D-mannitol, benzyl benzoate, ethanol, trisaminomethane, cholesterol, triethanolamine, sodium carbonate, sodium citrate, etc. Examples of the suspending agent include surfactants such as stearyl triethanolamine, sodium laurylsulfate, laurylaminopropionic acid, lecithin, benzalkonium chloride, benzetonium chloride, glycerin monostearate, etc.; hydrophilic polymers such as polyvinylalcohol, polyvinylpyrrolidone, sodium carboxymethyl cellulose, methyl cellulose, hydroxymethyl cellulose, hydroxyethyl cellulose, hydroxypropyl cellulose, etc.; etc. Examples of the isotonizing agent include sodium chloride, glycerin, D-mannitol, etc. Examples of the buffer include a buffer solution of phosphate, acetate, carbonate, citrate, etc. Examples of the soothing agent include benzylalcohol, etc. Examples of the preservative include paraoxybenzoic acid esters, chlorobutanol, benzylalcohol, phenethylalcohol, dehydroacetic acid, sorbic acid, etc. Examples of the

antioxidant include sulfites, ascorbic acid, etc.

The compound of the formula (I) of the present invention may be used in combination with other drug for the treatment or prevention of infectious disease of HIV (in particular, a pharmaceutical composition for the treatment or prevention of AIDS). In this case, these drugs can be formulated by mixing individually or simultaneously with pharmaceutically acceptable carriers, excipients, binders, diluents or the like, which can be administered non-orally as a pharmaceutical composition for the treatment or prevention of infectious disease of HIV. In the case of formulating these effective components individually, while the individually formulated agents can be administered in the form of their mixture prepared by using e.g. a diluent when administered, the individually formulated agents can also be administered separately or simultaneously or with time intervals to the one and same subject. A kit for administering the individually formulated effective components in the form of their mixture prepared by using e.g. a diluent when administered (e.g. a kit for injection which comprises two or more ampoules each comprising a powdery component and a diluent for mixing and dissolving two or more components when administered, etc.), etc. are also included by the pharmaceutical composition of the present invention.

Example of the other pharmaceutical agent for the treatment or prevention of infectious disease of HIV to be used in combination with the compound of the formula (I) of the present invention include nucleoside reverse transcriptases inhibitor such as zidovudine, didanosine, zalcitabine, lamivudine, stavudine, abacavir, adefovir, adefovir dipivoxil, fozivudine tidoxil, etc.; non-nucleoside reverse transcriptases inhibitor (including an agent having anti-oxidation activity such as immunocal, oltipraz, etc.) such as nevirapine, delavirdine, efavirenz, loviride, immunocal, oltipraz, tc.; protease inhibitors

such as saquinavir, ritonavir, indinavir, nelfinavir, amprenavir, palinavir, lasinavir, tc.; etc.

As the nucleoside reverse transcriptase inhibitor, zidovudine, didanosine, zalcitabine, lamivudine, stavudine, etc. are preferable; as the non-nucleoside reverse transcriptase inhibitor, nevirapine, delavirdine, etc. are preferable; and as the protease inhibitor, saquinavir, ritonavir, indinavir, nelfinavir, etc. are preferable.

10 The compound of the formula (I) can be produced in accordance with per se known methods, for example, the methods described below, the methods described in JP-A-73476/1996, or analogous methods thereto.

A salt of the compound of the formulas (I), (II), (III), (IV), (V), (I-1), (I-2) and (I-3) may form a salt with inorganic base, organic base, inorganic acid, organic acid, basic or acidic amino acid, etc., as long as such a salt does not interfere the reaction. Examples of the salt with the inorganic base include a salt with alkali metal (e.g. sodium, potassium, etc.), alkaline earth metal (e.g. calcium, magnesium, etc.), aluminum, ammonium, etc. Examples of the salt with the organic base include a salt with trimethylamine, triethylamine, pyridine, picoline, ethanolamine, diethanolamine, triethanolamine, dicyclohexylamine, N,N'-dibenzylethylenediamine, etc. Examples of the salt with the inorganic acid include a salt with hydrochloric acid, hydrobromic acid, nitric acid, sulfuric acid, phosphoric acid, etc. Examples of the salt with the organic acid include a salt with formic acid, acetic acid, trifluoroacetic acid, fumaric acid, oxalic acid, tartaric acid, maleic acid, citric acid, succinic acid, malic acid, methanesulfonic acid, benzenesulfonic acid, p-toluenesulfonic acid, etc. Examples of the salt with the basic amino acid include a salt with arginine, lysine, ornithine, etc. Examples of the salt with the acidic amino acid include a salt with aspartic acid, glutamic acid, etc.

In the following reaction steps, when the starting compounds have, as substituents, an amino group, a carboxyl group and/or hydroxy group, these groups may be protected by ordinary protective groups such as those generally employed in peptide chemistry, etc. After the reaction, if necessary, the protective groups may be removed to obtain the desired compound.

Examples of the amino-protective group include an optionally substituted formyl, an optionally substituted C₁₋₆ alkyl-carbonyl (e.g. methylcarbonyl, ethylcarbonyl, etc.), phenylcarbonyl, C₁₋₆ alkyloxy-carbonyl (e.g. methoxycarbonyl, ethoxycarbonyl, t-butoxycarbonyl, etc.), aryloxycarbonyl (e.g. phenoxycarbonyl, etc.), C₇₋₁₀ aralkyloxy-carbonyl (e.g. benzyloxycarbonyl, etc.), trityl, phthaloyl, etc. These protective groups may be substituted by 1 to 3 substituents such as halogen atom (e.g. fluorine, chlorine, bromine, iodine, etc.), C₁₋₆ alkyl-carbonyl (e.g. acetyl, propionyl, butyryl, etc.), nitro group, etc.

Examples of the carboxyl-protective group include an optionally substituted C₁₋₆ alkyl (e.g. methyl, ethyl, propyl, isopropyl, butyl, tert-butyl, etc.), phenyl, trityl, silyl, etc. These protective groups may be substituted by 1 to 3 substituents such as halogen atom (e.g. fluorine, chlorine, bromine, iodine, etc.), formyl, C₁₋₆ alkyl-carbonyl (e.g. acetyl, propionyl, butyryl, etc.), nitro group, etc.

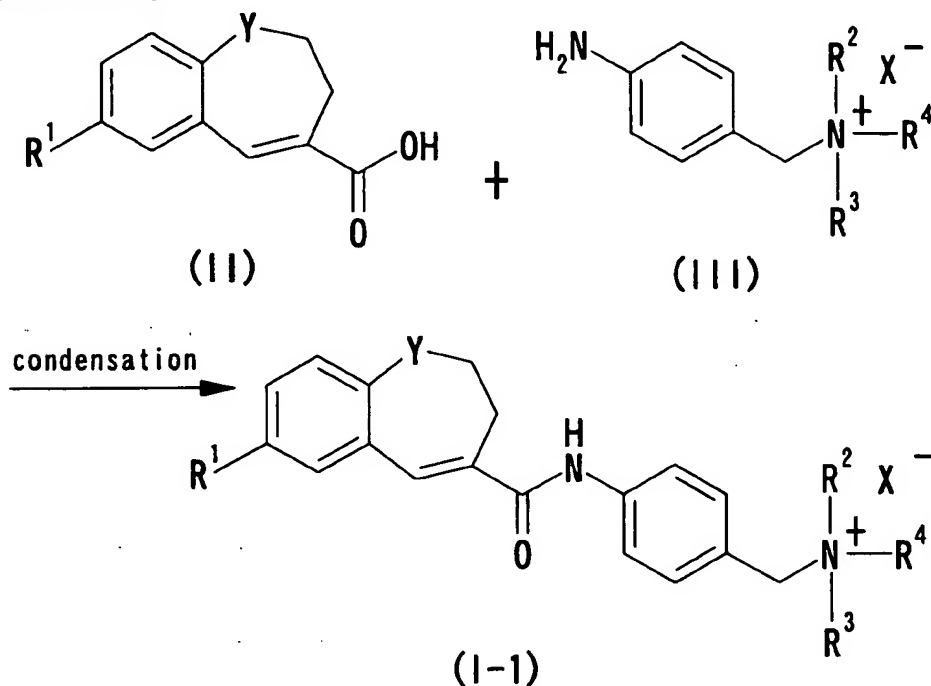
Examples of the hydroxy-protective group include an optionally substituted C₁₋₆ alkyl (e.g. methyl, ethyl, propyl, isopropyl, butyl, tert-butyl, etc.), phenyl, C₇₋₁₀ aralkyl (e.g. benzyl, etc.), formyl, C₁₋₆ alkyl-carbonyl (e.g. acetyl, propionyl, etc.), phenyloxycarbonyl, C₇₋₁₀ aralkyloxy-carbonyl (e.g. benzyloxycarbonyl, etc.), pyranyl, furanyl, silyl, etc. These protective groups may be substituted by 1 to 4 substituents such as halogen atom (e.g. fluorine, chlorine, bromine, iodine, etc.), C₁₋₆ alkyl, phenyl, C₇₋₁₀ aralkyl, nitro group, etc.

These protective group may be introduced or removed

by per se known methods (.g. a method described in Protective Groups in Organic Chemistry (J. F. W. McOmie et al.; Plenum Press Inc.) or the methods analogous thereto.

For example, employable method for removing the protective groups is a method using an acid, a base, reduction, ultraviolet ray, hydrazine, phenylhydrazine, sodium N-methyldithiocarbamate, tetrabutylammonium fluoride, palladium acetate, etc.

[Method A]



10

wherein X^- is an counter anion (e.g. an anion of a halogen atom (e.g. Cl^- , Br^- , I^- , etc.), etc.) and the other symbols are as defined above.

15

This production method is carried out by reacting the compound (II) with the aniline derivative (III) to obtain the anilide Compound (I-1).

The condensation reaction of the compounds (II) and (III) is carried out by usual methods for peptide synthesis.

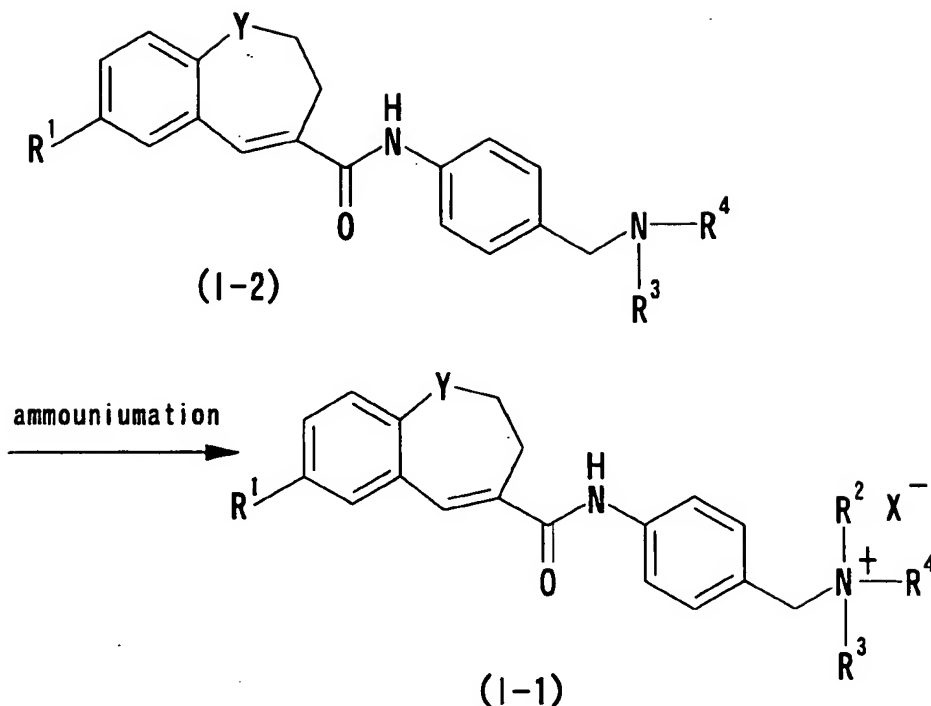
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Said methods for peptide synthesis are employed according

to optional known methods, for example, methods described in "Peptide Synthesis" written by M. Bodansky and M. A. Ondetti, Interscience, New York, 1966; "The Proteins", volume 2, written by F. M. Finn and K. Hofmann, H. Neurath and R. L. Hill edition, Academic Press Inc., New York, 1976; "peputido-gosei no kiso to jikken (Basis and Experiment of Peptide Synthesis)" written by Nobuo Izumiya et al., Maruzen K.K., 1985; etc., as well as azide method, chloride method, acid anhydride method, mixed acid anhydride method, DCC method, active ester method, method using Woodward reagent K, carbonyldiimidazole method, oxidation-reduction method, DCC/HONB method, etc. and in addition WSC method, method using diethyl cyanophosphate (DEPC), etc.

The condensation reaction can be carried out in a solvent. Examples of the solvents to be employed in the reaction include anhydrous or hydrous N,N-dimethylformamide (DMF), dimethylsulfoxide, pyridine, chloroform, dichloromethane, tetrahydrofuran, dioxane, acetonitrile, or a suitable mixture of these solvents. The reaction temperature is generally about -20°C to about 50°C, preferably about -10°C to about 30°C and the reaction time is generally about 1 to about 100 hours, preferably about 2 to about 40 hours.

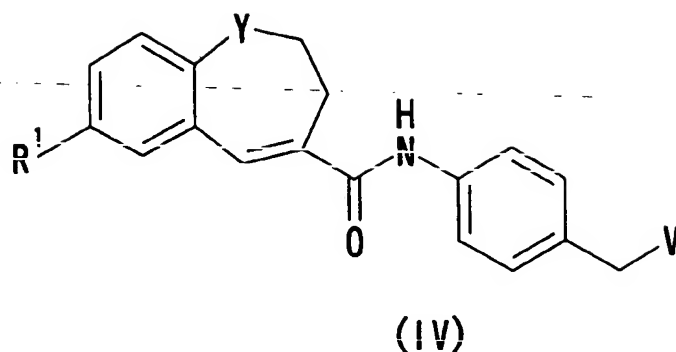
[Method B]



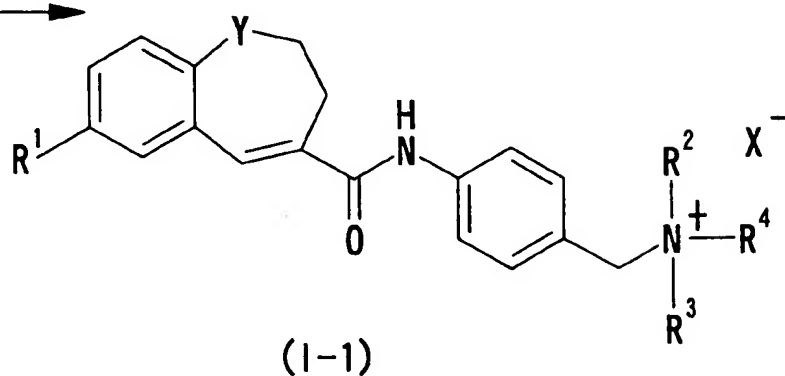
When Compound (I-2) has a tertiary amine residue, Compound (I-1) having an quaternary ammonium can be produced by reacting Compound (I-2) with halogenated alkyl.

- 5 Examples of a halogen atom include chlorine, bromine, iodine, etc. and usually about 1 to 5 moles of the halogenated alkyl (e.g. halogenated lower (C₁₋₆) alkyl, etc.) is used per mole of Compound (I-2). The reaction is carried out in an inert solvent such as toluene, benzene, xylene, dichloromethane,
- 10 chloroform, 1,2-dichloroethane, dimethylformamide, dimethylacetamide, etc., or a suitable mixture of these solvents. The reaction temperature is generally about 10°C to about 160°C, preferably about 20°C to about 120°C and the reaction time is generally about 1 hour to about 100 hours,
- 15 preferably about 2 hours to about 40 hours. This reaction is preferably carried out under inert gas (e.g. nitrogen, argon, etc.) atmosphere.

[Method C]



ammoniumation

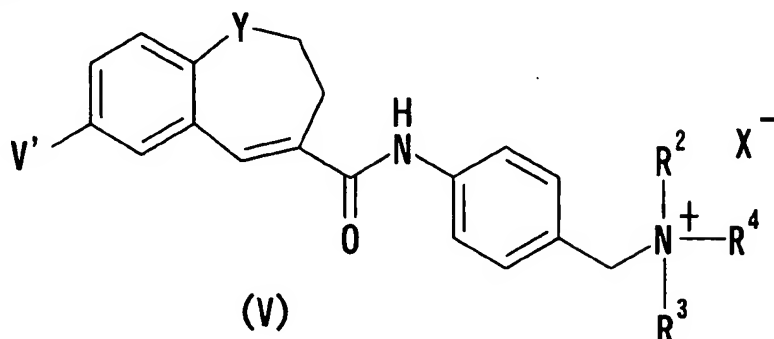


wherein V in the Compound (IV) is a halogen atom (chlorine, bromine, iodine, etc.), or a sulfonyloxy group (methanesulfonyloxy group, trifluoromethanesulfonyloxy group, benzenesulfonyloxy group, toluenesulfonyloxy group, etc.), and the other symbols are as defined above.

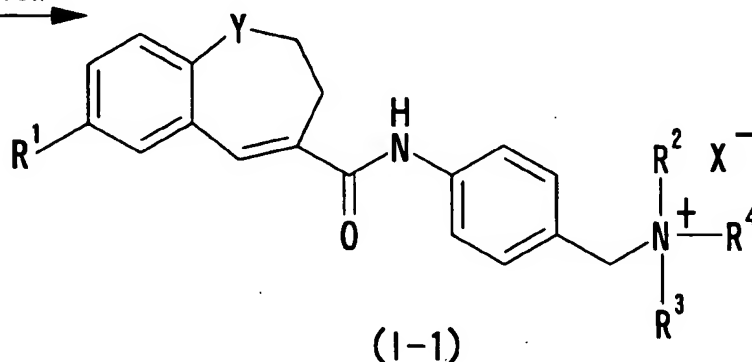
Compound (I-1) having a quaternary ammonium can be produced by reacting Compound (IV) and a tertiary amine. The reaction is carried out in an inert solvent such as toluene, benzene, xylene, dichloromethane, chloroform, 1,2-dichloroethane, dimethylformamide (DMF), dimethylacetamide, etc., or a suitable mixture of these solvents. Usually, about 1-3 moles of the tertiary amine is used per mole of Compound (IV). The reaction temperature is generally about 10°C to about 120°C, and the reaction time

is generally about 1 hour to about 40 hours. This reaction is preferably carried out under inert gas (e.g. nitrogen, argon, etc.) atmosphere.

[Method D]



Suzuki reaction



5

Compound (I-1) can be produced by subjecting Compound (V) wherein V' is a halogen atom (bromine, iodine, etc.) or a sulfonyloxy group (trifluoromethanesulfonyloxy group, etc.), and the other symbols are as defined above to, for example, Suzuki reaction [cross condensation reaction of aryl borate with e.g. aryl halide or aryloxytrifluoromethanesulfonate in the presence of palladium catalyst; A. Suzuki et al., Synth. Commun. 1981, 11, 513]. Usually, about 1-1.5 times moles of aryl borate is used per mole of

10

15 Compound (V).

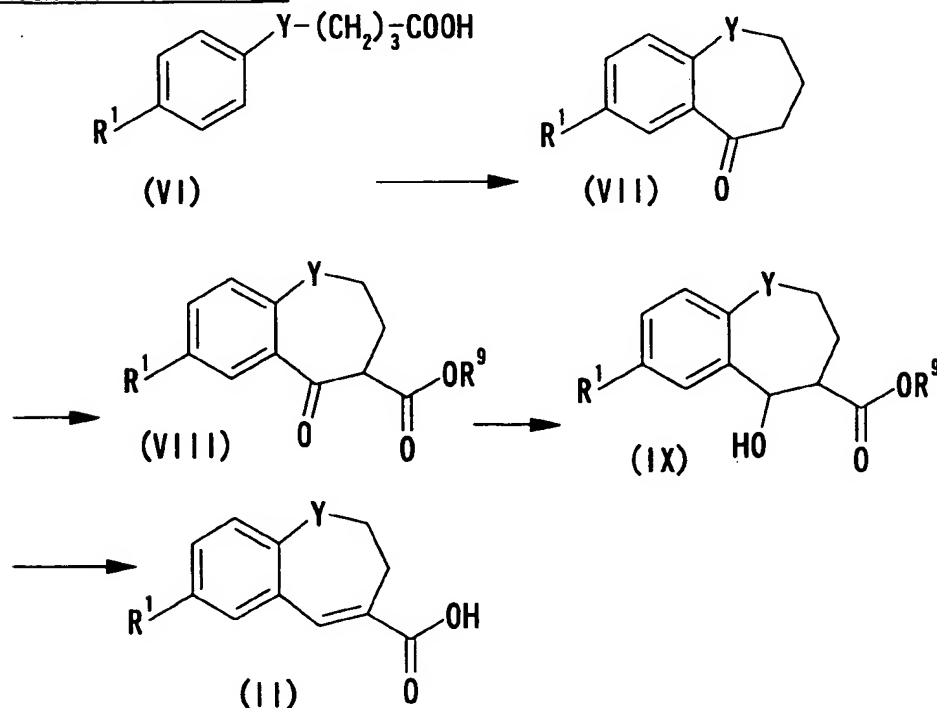
The thus obtained anilide derivative (I-1) can be isolated and purified by known separation and purification

methods such as concentration, concentration under reduced pressure, extraction, crystallization, recrystallization, solvent convert, chromatography, etc.

If necessary, the thus obtained compound (I-1) can be converted into the desired compound (I) wherein X is chlorine by per se known method (e.g. a method using ion exchange resin, exchange reaction in a solvent containing an excess amount of sodium chloride, etc.).

Compound (II) used as a starting material can be produced by a known method (e.g. method described in JP-A-73476/1996, etc.) or the methods analogous thereto. For example, Compound (II) can be produced by a method described in the following Reaction Scheme I, a method described in the following Reference Examples or the methods analogous thereto.

Reaction Scheme I

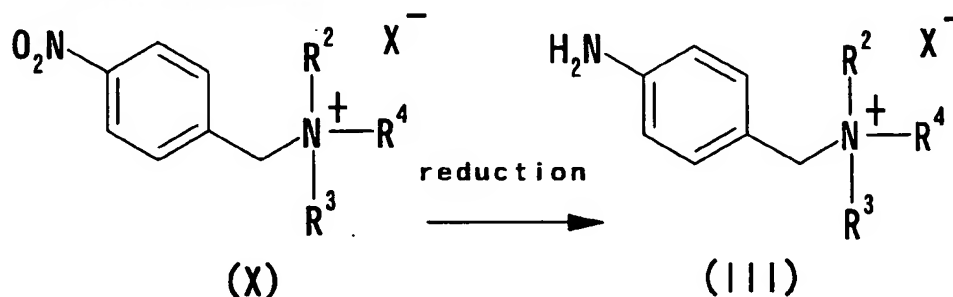


wherein R^1 is a C_{1-4} alkyl group, and the other symbols are as defined above.

In this reaction, the compound of the formula (VI) is heated with a polyphosphoric acid, or Compound (VI) is converted to acid chloride with thionyl chloride, oxalyl chloride, phosphorous oxychloride, phosphorous pentachloride, etc., followed by subjecting the resulting acid chloride to usual Friedel-Crafts reaction and cyclizing the same to produce Compound (VII). Compound (VII) is reacted with carbonate ester in the presence of a base to produce ketoester (VIII). Compound (VIII) is subjected to reduction with catalytic hydrogenation or sodium boron hydride, etc. to produce Compound (IX). Compound (IX) is subjected to dehydration and ester hydrolysis by per se known methods to produce unsaturated carboxylic acid (II).

Compound (III) can be produced by a known method (e.g. method described in JP-A-73476/1996, etc.) or the methods analogous thereto. For example, Compound (III) can be produced by a method described in the following Reaction Scheme II, a method described in the following Reference Examples or the methods analogous thereto.

Reaction Scheme II



The reduction of Compound (X) can be carried out by per se known methods, for example, reduction with metal, reduction with metal hydride, reduction with metal hydride complex compound, reduction with diborane or substituted borane, catalytic hydrogenation, etc. That is, this reaction is carried out by treating Compound (X) with reduction agent. Examples of the reduction agent include

metal such as reduced iron, zinc powder, etc.; alkali metal boron hydride (e.g. sodium boron hydride, lithium boron hydride, etc.); metal hydride complex compound such as aluminum lithium hydride, etc.; metal hydride such as sodium hydride etc.; organic tin compound (triphenyltin hydride, etc.), metal complex compound and metal salt such as nickel compound, zinc compound etc.; catalytic reduction agent using hydrogen and transit metal catalyst such as palladium, platinum, rhodium, etc.; diborane; etc. Among others, as the reduction agent, catalytic reduction agent using hydrogen and transit metal catalyst such as palladium, platinum, rhodium, etc.; reduced iron, etc. are preferable. The reaction is carried out in a solvent which does not inhibit the reaction. Examples of the solvent include benzene, toluene, xylene, chloroform, carbon tetrachloride, dichloromethane, 1,2-dichloroethane, 1,1,2,2-tetrachloroethane, diethylether, tetrahydrofuran, dioxane, methanol, ethanol, propanol, isopropanol, 2-methoxyethanol, N,N-dimethylformamide, acetic acid, or a suitable mixture of these solvents, etc. The solvent is appropriately selected depending on kind of the reduction agent. The reaction temperature is generally about -20°C to about 150°C, preferably about 0°C to about 100°C, and the reaction time is generally about 1 to about 24 hours.

The resulting Compound (III) can be separated and purified with know separation and purification methods such as concentration, concentration under reduced pressure, extraction, crystallization, was recrystallized with, solvent conversion, chromatography, etc.

The compound of the formula (I) of the present invention has potent CCR5 antagonistic activity and therefore can be used for the treatment or prevention of various infectious diseases of HIV, for example, AIDS in human. The compound of the formula (I) of the present invention is low toxic and safely used as CCR5 antagonist for the treatment or prevention of AIDS and also for th prevention of the

progression of AIDS. The compound of the formula (I) of the present invention has superior solubility and absorbability, and therefore it can be advantageously used as injection.

The dose per day of the compound of the formula (I) varies depending on the condition and body weight of a patient, administration route, etc. Typical daily dose per adult patient (body weight: 50Kg) for subcutaneous administration is about 1-200mg, preferably about 1-100mg, more preferably about 2-50mg, and in particular about 5-30mg, as active ingredient [the compound of the formula (I)] and the compound of the formula (I) is administered once or 2-3 times par day.

When the compound of the formula (I) is used in combination with a reverse transcriptase inhibitor and/or a protease inhibitor, the dose of the reverse transcriptase inhibitor or the protease inhibitor ranges, for example, from about 1/200-1/2 or more of usual dose to about 2-3 times or less of usual dose. In case that two or more drugs are used in combination, each dose of the drugs is appropriately adjusted if one drug affects metabolism of the other drug, while each dose of the drugs when they are used in combination is generally the same as the dose when they are used alone.

Typical daily dose of the reverse transcriptase inhibitor and the protease inhibitor is as follows:

zidovudine	: 100mg
didanosine	: 125-200mg
zalcitabine	: 0.75mg
lamivudine	: 150mg
stavudine	: 30-40mg
saquinavir	: 600mg
ritonavir	: 600mg
indinavir	: 800mg
nelfinavir	: 750mg

In case of combination use of the compound of the formula (I) with a reverse transcriptase inhibitor and/or

a protease inhibitor preferred embodiments are shown below.

- ① A drug containing about 1-200mg of the compound of the formula (I) and a drug containing about 50-200mg of zidovudine to one adult patient (body weight: 50Kg) are administered. Each of the drugs may be administered to the one and the same subject simultaneously or with time intervals of 12 hours or less.
- ② A drug containing about 1-200mg of the compound of the formula (I) and a drug containing about 300-1200mg of saquinavir to one adult patient (body weight: 50Kg) are administered. Each of the drugs may be administered to the one and the same subject simultaneously or with time intervals of 12 hours or less.

15

Best Mode for Carrying out the Invention

The present invention is hereinafter described in more detail by means of the following Test Example, Reference Example and Working Example, which are mere examples of the present invention and are not construed as limitative to the present invention.

The following gene manipulation is carried out in accordance with methods described in textbook (Maniatis et al., Molecular Cloning, Cold Spring Harbor Laboratory, 1989) or protocol attached to reagents.

25

Test Example

(1) Cloning of human CCR5 chemokine receptor

Cloning of CCR5 gene was carried out by PCR (polymerase chain reaction) from human spleen cDNA. With using 0.5ng of spleen cDNA (Toyobo, QUICK-Clone cDNA) as template, PCR was performed in DNA Thermal Cycler 480 (Perkin-Elmer) (reaction conditions: 30 cycles of 95°C for 1 minute, 60°C for 1 minute, and 75°C for 5 minutes) by adding primer set,

35 5'-CAGGATCCGATGGATTATCAAGTGTCAAGTCCAA-3' (25pmol) and
5'-TCTAGATCACAAGCCCACAGATATTTCTGCTCC-3' (25pmol),

which were designed referring to nucleotide sequence of CCR5 gene reported by Samson et al. (Biochemistry, 35(11), 3362-3367 (1996)) and by using TaKaRa EX Taq (Takara Shuzo). The resultant PCR product was subjected to agarose gel electrophoresis to collect about 1.0kb DNA fragment, which was subjected to Original TA Cloning Kit (Funakoshi) to carry out cloning of CCR5 gene.

(2) Preparation of plasmid for expression of human CCR5

The plasmid obtained in the above (1) was digested with restriction enzymes XbaI (Takara Shuzo) and BamHI (Takara Shuzo) and subjected to agarose gel electrophoresis to collect about 1.0kb DNA fragment. The DNA fragment was mixed with plasmid pcDNA3.1 (Funakoshi) for expression in animal cells, said plasmid being digested with XbaI and BamHI, and they were ligated with DNA Ligation Kit Ver.2 (Takara Shuzo). The resulting plasmid was subjected to transformation of competent cell of E. coli JM109 (Takara Shuzo) to obtain plasmid pCKR5.

(3) Introduction of plasmid for expression of human CCR5 into CHO-K1 cell and Expression of plasmid for expression of human CCR5 in CHO-K1 cell

CHO-K1 cells were grown in 750ml of tissue culture flask (Becton Dickinson) using Ham's F12 medium (Nihon Pharmaceutical) containing 10% fetal calf serum (Life Tech Oriental) and took off with 0.5g/L trypsin-0.2g/L EDTA (Life Tech Oriental). The cells were washed with PBS (Life Tech Oriental), centrifuged (1000rpm, 5 minutes), and suspended in PBS. With using Gene Pulser (Bio-Rad Laboratories), DNA was introduced into the cells under the conditions shown below. That is, to the cuvette of 0.4cm gap were added 8×10^6 cells and 10 μ g of plasmid pCKR5 for expression of human CCR5, and electroporation was carried out under 0.25kV of voltage and 960 μ F of capacitance.

The cells were transferred into Ham's F12 medium (Nihon Pharmaceutical) containing 10% fetal calf serum, and cultivated for 24 hours. The cells were again took off and

centrifuged, and suspended in Ham's F12 medium (Nihon Pharmaceutical) containing 10% fetal calf serum and 500 $\mu\text{g/ml}$ of geneticin (Life Tech Oriental). The suspension was diluted to give 10^4 cells/ml of the suspension, which
5 was inoculated on 96 well plate (Becton Dickinson) to give geneticin resistant cells. The resulting geneticin resistant cells were cultivated in 96 well plate (Becton Dickinson), and cells expressing CCR5 were selected from the geneticin resistant cells. That is, in assay buffer
10 (Ham's F12 medium containing 0.5% BSA and 20mM HEPES (Wako Pure Chemical, pH7)) to which was added 200pM of [^{125}I]-RANTES (Amersham) as ligand, binding reaction was carried out at room temperature for 40 minutes, and the buffer was washed with cooled PBS. To the buffer was added 50 μl /well of 1M
15 NaOH, and the mixture was stirred. Radioactivity was determined with γ -counter to select CHO/CCR5 cells which specifically bind to the ligand.

(4) Evaluation of Test Compounds based on CCR5 antagonistic activity

20 The CHO/CCR5 were inoculated on 96 well microplate (5×10^4 cells/well) and cultivated for 24 hours. The medium was removed by means of suction, and to each well was added assay buffer containing Test Compound ($1 \mu\text{M}$) and then 100pM of [^{125}I]-RANTES (Amersham) as ligand. Binding assay was
25 carried out at room temperature for 30 minutes, and assay buffer was removed by means of suction. Each well was washed twice with cooled PBS, and 200 μl of Microscint-20 (Packard Instrument, Inc.) was added to each well. Radio-activity was determined with TopCount Micro Scintillation Counter
30 (Packard Instrument, Inc.).

According to the method described above, inhibition rate of Test Compound (whose number is referred to in the following Examples) to CCR5 binding.

The results are shown in Table 1.

35

Table 1

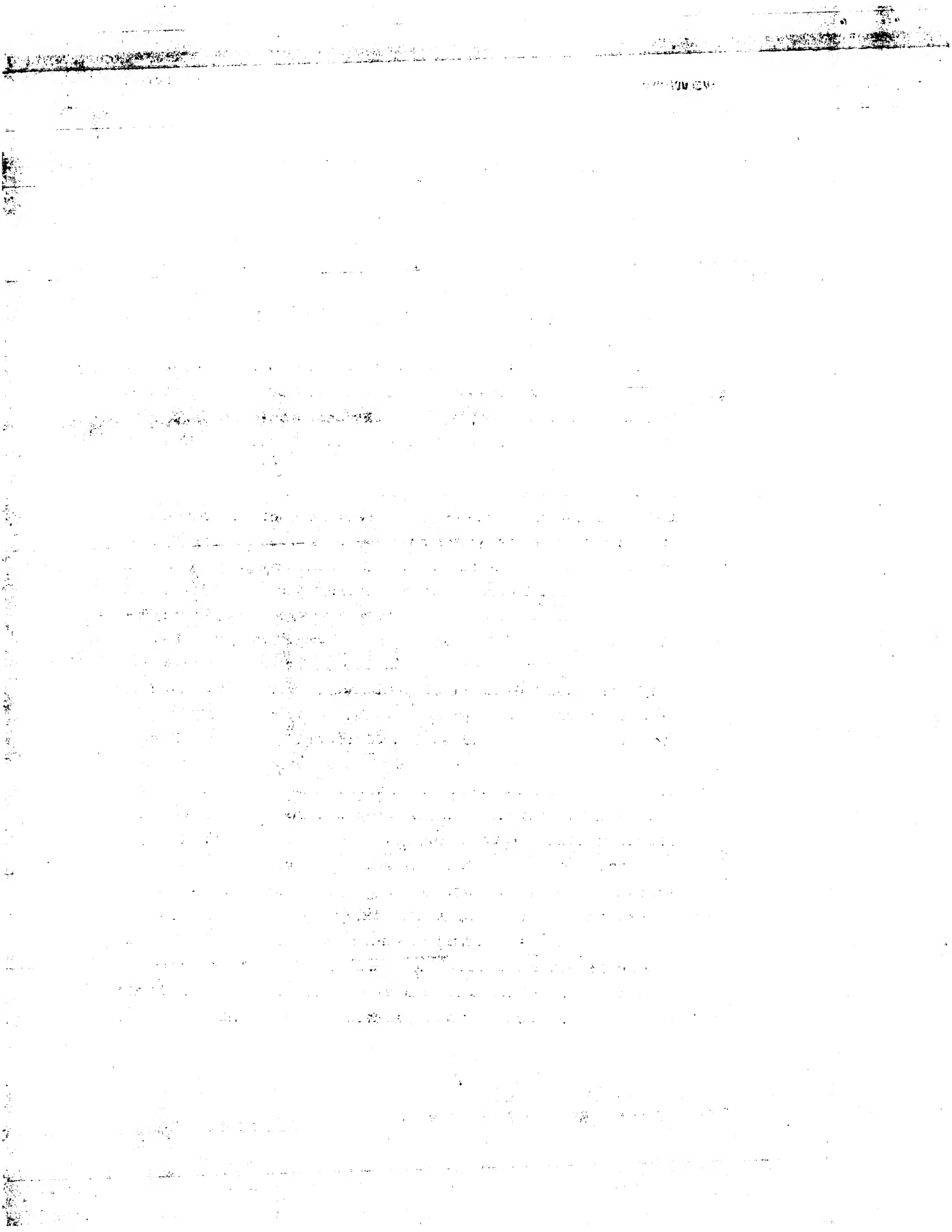
	Compound Number	Inhibition Rate (%)
	1	99
	2	96
5	3	96

(5) Inhibitory effect on HIV-1 infection to MAGI-CCR5 cell

The plasmid where β -galactosidase gene was ligated downstream of HIV-1 LTR was introduced into CD4 positive HeLa cell, to which human CCR5 was further introduced to obtain transformant MAGI-CCR5. By using said transformant MAGI-CCR5, degree of HIV-1 infection was calculated from β -galactosidase activity (blue color due to decomposition of 5-bromo-4-chloro-3-indolyl- β -D-galactopyranoside).

Specifically, MAGI-CCR5 cells were suspended in DMEM medium containing 10% serum to prepare 5×10^4 cells/ml suspension. To each well of 96 well plate was inoculated $200 \mu\text{l}$ of the suspension, and the cells were cultivated at 37°C overnight. The medium was removed by means of suction, and to the residue was added $100 \mu\text{l}$ of the above medium containing $0.064 \mu\text{M}$ of Test Compound and $100 \mu\text{l}$ of the above medium containing 300PFU of HIV-1 BA-L cells. The cells were cultivated at 37°C for 2 days. The medium was removed by means of suction.

To the residue was added $200 \mu\text{l}$ of cell fixative (PBS containing 1% formaldehyde and 0.2% glutaraldehyde), and the mixture was allowed to stand at room temperature for 5 minutes and washed twice with PBS. To the mixture was added $100 \mu\text{l}$ of staining solution (PBS containing $4 \mu\text{M}$ potassium ferrocyanide, $4 \mu\text{M}$ potassium ferricyanide, $2 \mu\text{M}$ MgCl_2 and 0.4mg/ml X-gal), and the mixture was allowed to stand at 37°C for 50 minutes and washed twice with PBS. The number of blue cells was counted by microscope and defined as the number of cells infected with HIV-1. According to this method, inhibition rate on HIV-1 infection was determined and found that Compound No. 1 shows 100% inhibition on HIV-1 infection.



(6) Inhibitory effect on HIV-1 infection to human PBMC

From normal person human peripheral blood mononuclear cells (PBMC) were separated, and the cells were stimulated with 10 μ g/ml of PHA (Phytohemagglutinin) and 20U/ml of interleukin-2 (IL-2) for 3 days. The cells were suspended in RPMI-1640 medium containing 20% serum to prepare 1×10^6 /ml suspension. To the suspension were infected HIV-1 BA-L cells (20ng as an amount of p24 antigen), and viruses were absorbed at 37°C for 2 hours. The cells were washed and suspended in RPMI-1640 medium containing 20% serum and IL-2 20U/ml to prepare 1×10^5 /ml suspension. To the PBMC suspension was added the same amount of a solution which contains 0.032 μ M of Test Compound, and the cells were cultivated at 37°C in carbon dioxide gas incubator. The amount of p24 antigen in supernatant of the cultivated medium was determined by enzyme-linked immunosorbent assay (ELISA) and defined as degree of HIV-1 infection. According to this method, inhibition rate on HIV-1 infection was determined and found that Compound No. 1 shows 74% inhibition on HIV-1 infection.

The pharmaceutical composition for antagonizing CCR5 (e.g. a medicament for the treatment or prevention of infectious disease of HIV, a medicament for the treatment or prevention of AIDS, etc.) comprising the compound of the formula (I) of the present invention, as an active ingredient, can be prepared, for example, by the following prescriptions:

1. Injection

A mixture of Compound 1 (500mg), mannitol (1000mg) and polysorbate 80 (100mg) is dissolved in distilled water (10ml), and to the solution is added distilled water to make the whole volume 20ml. The solution is filtered under sterile conditions. Each 2ml of the solution is filled into a vial for injection under sterile conditions.

Reference Example 1

To a solution of 4-nitrobenzylalcohol (50 g, 0.326 mol) in ethyl acetate (EtOAc) (200 ml) were added 3,4-dihydropyran (35.7 ml, 0.392 mol) and CSA (camphor sulfonic acid) (379 mg, 1.63 mmol) under stirring at room temperature, and the mixture was stirred at room temperature for 1 hour. After the reaction completed, the reaction mixture was neutralized with saturated NaHCO₃ solution and separated ethyl acetate layer was dried with MgSO₄ and concentrated under reduced pressure. The residue was purified with silica gel column chromatography to give 4-(2-tetrahydropyranyloxymethyl)nitrobenzene (74.5 g, 96%) as syrup.

¹H-NMR (200 MHz, CDCl₃) δ: 1.55-2.05 (6H, m), 3.51-3.62 (1H, m), 3.83-3.94 (1H, m), 4.61 (1H, d, J=13.6Hz), 4.74 (1H, t, J=3.2Hz), 4.93 (1H, d, J=13.4Hz), 7.51-7.56 (2H, d, J=8.8Hz), 8.18-8.24 (2H, m).

Reference Example 2

To a solution of 4-(2-tetrahydropyranyloxymethyl)-nitrobenzene (59.7 g, 0.256 mol) in ethanol (EtOH) (300 ml) was added under nitrogen atmosphere at room temperature 10% Pd/C (5.97 g), and catalytic hydrogenation was carried out. The mixture was stirred at room temperature for 24 hours. After the reaction completed, the catalyst was filtered off, and the organic layer was concentrated under reduced pressure. The residue was purified with silica gel column chromatography to give 4-(2-tetrahydropyranyloxymethyl)-aniline (39.7 g, 76%) as syrup.

¹H-NMR (200 MHz, CDCl₃) δ: 1.45-1.95 (6H, m), 3.00-3.60 (3H, br m), 3.87-4.14 (1H, m), 4.39 (1H, d, J=11.4Hz), 4.68 (1H, d, J=11.4Hz), 4.71 (1H, m), 6.65-6.69 (2H, m), 7.15-7.19 (2H, m).

Reference Example 3

To a solution of 2-(4-methylphenyl)-6,7-dihydro-5H-benzocycloheptene-8-carboxylic acid (35.0 g, 0.126 mol) in tetrahydrofuran (THF) (280 ml) were added (COCl)₂ (21.9 ml, 0.251 mol) and DMF (0.7 ml) at 0°C. Under nitrogen

atmosphere, the mixture was stirred at room temperature for 4 hours. After the reaction completed, the solvent was evaporated, and to the residue was added THF (315 ml). To a solution of the acid chloride was added a solution of 4-(2-tetrahydropyranyloxymethyl)aniline (28.1 g, 0.138 mol) and triethylamine (Et₃N) (26.3 ml, 0.189 mol) in THF (105 ml) at 0°C, and the mixture was stirred under nitrogen atmosphere, at room temperature for 2 hours. After the reaction completed, to the mixture was added water, and the mixture was extracted with ethyl acetate. The organic layer was washed with saturated NaCl solution and dried with MgSO₄. The solvent was evaporated and the residue was dissolved in methanol (MeOH) (470 ml). To the mixture was dropwise added 6N HCl (5.9 ml) at room temperature, and the mixture was stirred for 1 hour. After the reaction completed, the mixture was neutralized with saturated NaHCO₃ solution, and the solvent was removed. The residue was washed with water and then acetone/isopropylether (10:1; 60 ml), and the resulting precipitate was filtered, which was dissolved in THF. The mixture was dried with MgSO₄, and the solvent was evaporated. The resulting powder was washed twice with hexane:ethyl acetate (10:1; 50 ml) to give N-(4-hydroxymethylphenyl)-2-(4-methylphenyl)-6,7-dihydro-5H-benzocycloheptene-8-carboxamide (26.8 g, 56%) as white powder.

¹H-NMR (200 MHz, CDCl₃) δ: 2.10-2.22 (2H, m), 2.39 (3H, s), 2.71 (2H, br t, J=6.4), 2.84-2.91 (2H, m), 4.67 (2H, s), 7.20-7.26 (2H, m), 7.33-7.51 (7H, m), 7.61 (2H, d, J=8.4), 7.71 (1H, br s).

Reference Example 4

To a solution of N-(4-hydroxymethylphenyl)-2-(4-methylphenyl)-6,7-dihydro-5H-benzocycloheptene-8-carboxamide (10.0 g, 26.1 mmol) and pyridine (0.1 ml) in chloroform (150 ml) was dropwise added a solution of thionyl chloride (3.4 ml, 39.2 mmol) in chloroform (90 ml), and the mixture was stirred under nitrogen atmosphere at room

temperature for 17 hours. After the reaction complet d,
water was added to the mixture, and th mixtur was extracted
with chloroform. The organic layer was washed with
saturated sodium chloride solution and dried with anhydrous
5 magnesium sulfate. The solvent was evaporated, and the
resulting powder was washed with hexane to give N-(4-
chloromethylphenyl)-2-(4-methylphenyl)-6,7-dihydro-5H-
benzocycloheptene-8-carboxamide (10.2 g, 97%) as
colorless powder.

10 $^1\text{H-NMR}$ (200 MHz, CDCl_3) δ : 2.05-2.21 (2H, m), 2.40 (3H, s),
2.71 (2H, br t, $J=6.4$), 2.84-2.91 (2H, m), 4.58 (2H, s),
7.20-7.27 (2H, m), 7.35-7.52 (7H, m), 7.59-7.65 (2H, m),
7.71 (1H, br s).

Anal. for $\text{C}_{26}\text{H}_{24}\text{NOCl}\cdot 0.25\text{H}_2\text{O}$:

15 Calcd: C; 76.83, H; 6.08, N; 3.45.

Found: C; 76.55, H; 6.00, N; 3.53.

Reference Example 5

To a solution of tetrahydro-4H-pyran-4-one (60 g, 0.6
mol) and water (5 ml) in DMF (70 ml, 0.90 mol) was added
20 formic acid (46 ml, 1.2 mol), and the mixture was stirred
at 140°C for 23 hours. After the reaction completed, reflux
apparatus was changed to evaporation apparatus, crude amine
was obtained by evaporation (74.6 g).

b.p. $117 - 123^\circ\text{C}$ (27 mm).

25 To an aqueous solution (100 ml) of the crude amine (30
g) was dropwise added 6N HCl (5 drops), and the mixture was
washed twice with dichloromethane. The aqueous layer was
adjusted to pH 11 with sodium hydroxide. To the mixture was
added NaCl, and the mixture was extracted with
30 dichloromethane three times. The organic layer was dried
with potassium carbonate, and the solvent was evaporated.
The residue was purified with evaporation to give N,N-
dimethyl-N-tetrahydropyran-4-ylamine (10.4 g, 29%) as
colorless oil.

35 b.p. $75-82^\circ\text{C}$ (29 mm).

$^1\text{H-NMR}$ (200 MHz, CDCl_3) δ : 1.40-1.82 (4H, m), 2.28 (6H, s),

2.25-2.40 (1H, m), 3.37 (2H, ddd, J=11.8, 11.8 and 2.2),
3.97-4.05 (2H, m).

Reference Example 6

To a suspension of 7-(4-methylphenyl)-2,3-dihydro-
5 1-benzoxepine-4-carboxylic acid (0.6 g, 2.1 mmol) in
tetrahydrofuran (10 ml) were added oxalyl chloride (0.33
ml, 4.3 mmol) and N,N-dimethylformamide (1 drop) at 0°C, and
the mixture was stirred at room temperature for 2.5 hours.
The solvent was evaporated, and the residue was dissolved
10 in tetrahydrofuran (6 ml). To the mixture was dropwise
added 4-(t-butyldimethylsilyloxymethyl)aniline (0.56 g,
2.4 mmol) and triethylamine (0.36 ml, 2.6 mmol) in
tetrahydrofuran (2 ml) at 0°C, and the mixture was stirred
at room temperature for 16 hours. To the reaction mixture
15 was added water, and the mixture was extracted with ethyl
acetate. The extract was washed with saturated sodium
chloride solution and dried with magnesium sulfate. The
solvent was evaporated, and the residue was subjected to
silica gel column chromatography. Crude amide (1.1 g) was
20 obtained from fractions of hexane:ethyl acetate=5:1. This
product was dissolved in acetone (8 ml), and to the mixture
was dropwise added 6N hydrochloric acid. The mixture was
stirred for 1 hour. To the mixture were added 1% sodium
hydrogen carbonate (100 ml) and diisopropylether (100 ml),
25 and precipitate was filtered, which were dissolved in
acetone. The mixture was dried with magnesium sulfate, and
the solvent was evaporated. The resulting powder was
recrystallized from acetone-diisopropyl-ether to give
N-(4-hydroxymethylphenyl)-7-(4-methylphenyl)-2,3-
30 dihydro-1-benzoxepine-4-carboxamide (0.87 g) as colorless
crystals.

¹H-NMR (CDCl₃) δ: 2.39 (3H, s), 3.08 (2H, br t, J=4.4), 4.36
(2H, t, J=4.4), 4.68 (2H, s), 7.06 (2H, d, J=8.4), 7.18-7.61
(10H, m), 7.24 (2H, d, J=8.4).

35 Anal. for C₂₅H₂₃NO₃:

Calcd: C; 77.90, H; 6.01, N; 3.63.

Found: C; 77.91, H; 6.10, N; 3.55.

Referenc Example 7

To a solution of N-(4-hydroxymethylphenyl)-7-(4-methylphenyl)-2,3-dihydro-1-benzoxepine-4-carboxamide
5 (412 mg, 1.07 mmol) and pyridine (1 drop) in chloroform (5 ml) was dropwise added thionyl chloride (0.14 ml, 1.61 mmol), and the mixture was stirred for 2 hours. The mixture was diluted with water and extracted with chloroform. The extract was washed with saturated sodium chloride solution
10 and dried with magnesium sulfate. The solvent was evaporated, and the resulting powder was washed with hexane-ethyl acetate (1:1) to give N-(4-chloromethylphenyl)-7-(4-methylphenyl)-2,3-dihydro-1-benzoxepine-4-carboxamide (380 mg, 88%) as colorless powder.

15 m.p. 164°C

¹H-NMR (CDCl₃) δ: 3.29 (3H, s), 3.07 (2H, t, J=4.8), 4.36 (2H, t, J=4.8), 4.59 (2H, s), 7.05 (1H, d, J=8.2), 7.22-7.26 (2H, m), 7.36-7.52 (6H, m), 7.57-7.62 (3H, m).

Anal. for C₂₅H₂₂NO₂Cl:

20 Calcd: C; 74.34, H; 5.49, N; 3.47.

Found: C; 74.00, H; 5.42, N; 3.29.

Reference Example 8

To a suspension of 1,4-cyclohexanedione monoethylene-ketal (3.82 g, 24.6 mmol) and dimethylamine hydrochloride
25 (2.00 g, 24.6 mmol) in 1,2-dichloroethane (50 ml) were dropwise added triethylamine (4.2 ml, 29.6 mmol) and DBU (1,8-diazabicyclo-[5.4.0]-7-undecene) (4.4 ml), and the mixture was stirred for 10 minutes. To the mixture was added triacetoxyborohydride (7.68 g, 34.4 mmol), and the mixture
30 was stirred for 4.5 hours. Precipitate was filtered off, and the filtrate was concentrated to give crude product (6.34 g), which was dissolved in water (10 ml). To the mixture was dropwise added concentrated hydro-chloric acid (6 ml), and the mixture was stirred for 48 hours. The reaction
35 mixture was diluted with water and washed twice with ether. The aqueous layer was made basic with sodium hydroxide and

extracted with ether twice. The extract was washed with saturated sodium chloride solution, dried with potassium carbonate and purified by evaporation to give 4-dimethylaminocyclohexanone (0.59 g, 17%).

5 b.p. 142-145°C

¹H-NMR (CDCl₃) δ: 1.69-2.13 (4H, m), 2.32 (6H, s), 2.20-2.41 (2H, m), 2.44-2.64 (3H, m).

Reference Example 9

To a solution of 7-(4-ethoxyphenyl)-2,3-dihydro-1-benzoxepine-4-carboxylic acid (2.38 g) in THF (50 ml) were added oxalyl chloride (1.4 ml) and DMF (2 drops) at room temperature, and the mixture was stirred for 1 hour. Under reduced pressure, the solvent was evaporated, and the residue was dissolved in THF (50 ml). To the mixture was dropwise added a solution of triethylamine (2.1 ml) and 4-aminobenzyloxy-tert-butyldimethylsilane (2.00 g) in THF (10 ml) at 0°C, and the mixture was stirred at room temperature for 18 hours. To the reaction mixture was added water, and the mixture was extracted with ethyl acetate. The organic layer was washed with saturated sodium chloride solution, dried with magnesium sulfate and concentrated under reduced pressure. The residue was separated and purified with column chromatography (ethyl acetate/hexane = 1:4) to give pale yellow crystals (3.99 g), which were dissolved in acetone (50 ml). To the mixture was added 6N hydrochloric acid (1.3 ml) at room temperature, and the mixture was stirred for 1 hour. To the reaction mixture were added 5% sodium hydrogen carbonate solution (15 ml) and diisopropylether (100 ml). Precipitate was collected by filtration and washed with water and diisopropylether. The resulting solid was dissolved in THF, dried with magnesium sulfate and concentrated under reduced pressure to give crystals, which were recrystallized from THF to give 7-(4-ethoxyphenyl)-N-(4-hydroxymethylphenyl)-2,3-dihydro-1-benzoxepine-4-carboxamide (2.65 g) as colorless crystals.

m.p. 208-210 °C

¹H-NMR (200MHz, DMSO-d₆) δ: 1.35 (3H, t, J=7.0 Hz), 2.93-3.03 (2H, m), 4.06 (2H, q, J=7.0 Hz), 4.45 (2H, br s), 5.01-5.18 (1H, m), 6.98-7.05 (3H, m), 7.25-7.34 (3H, m),
5 7.49-7.71 (6H, m), 9.92 (1H, s).

IR (KBr) ν: 3363, 3290, 1659, 1612, 1525, 1493, 1242, 1227, 825 cm⁻¹

Anal. for C₂₆H₂₃NO₄

Calcd: C, 75.16 ; H, 6.06 ; N, 3.37

10 Found: C, 75.16 ; H, 6.08 ; N, 3.31.

Reference Example 10

To a suspension of 7-(4-ethoxyphenyl)-N-(4-hydroxy-methylphenyl)-2,3-dihydro-1-benzoxepine-4-carboxamide (2.55 g) and pyridine (2 drops) in chloroform (50 ml) was
15 added thionyl chloride (0.8 ml) at room temperature, and the mixture was stirred for 20 hours. To the reaction mixture was added water and then THF, and the mixture was extracted with ethyl acetate. The organic layer was washed with saturated sodium chloride solution, dried with
20 magnesium sulfate and concentrated under reduced pressure to give solid, which was dissolved in THF and ethyl acetate. The mixture was concentrated under reduced pressure to give crystals, which were collected by filtration and washed with diisopropylether to give N-(4-chloromethylphenyl)-7-(4-
25 ethoxyphenyl)-2,3-dihydro-1-benzoxepine-4-carboxamide (2.42 g) as colorless crystals.

m.p. 187-189 °C

¹H-NMR (200MHz, DMSO-d₆) δ: 1.35 (3H, t, J=7.0 Hz), 2.93-3.04 (2H, m), 4.06 (2H, q, J=7.0 Hz), 4.23-4.34 (2H, m),
30 4.74 (2H, s), 6.98-7.06 (3H, m), 7.35-7.42 (3H, m), 7.52 (1H, dd, J=8.4, 2.2 Hz), 7.59 (2H, d, J=8.8 Hz), 7.70-7.74 (3H, m), 10.04 (1H, s).

IR (KBr) ν: 3400, 1659, 1610, 1525, 1493, 1242, 1047, 822 cm⁻¹

35 Anal. for C₂₆H₂₄NO₃Cl

Calcd: C, 71.97 ; H, 5.57 ; N, 3.23

Found: C, 71.96 ; H, 5.54 ; N, 3.04.

Reference Example 11

To solution of 7-(4-ethoxyphenyl)-N-[4-[N-methyl-N-(tetrahydropyran-4-yl)aminomethyl]phenyl]-2,3-dihydro-1-benzoxepine-4-carboxamide (111 mg) in DMF (5 ml) was added methyl iodide (0.04 ml) at room temperature, and the mixture was stirred for 8 hours. Under reduced pressure, the mixture was concentrated, and to the residue was added ethyl acetate to precipitate solid, which was collected by filtration and recrystallized from ethanol-ethyl acetate to give dimethyl-[4-N-[7-(4-ethoxyphenyl)-2,3-dihydro-1-benzoxepin-4-carbonyl]aminobenzyl]-4-tetrahydropyranylammonium iodide (97 mg) as pale yellow crystals. m.p. 152-158 °C

¹H-NMR (200MHz, CDCl₃) δ: 1.41 (3H, t, J=7.0 Hz), 1.68-1.98 (2H, m), 2.10-2.26 (2H, m), 2.94 (6H, s), 2.98-3.08 (2H, m), 3.35-3.59 (3H, m), 3.96-4.16 (2H, m), 4.03 (2H, q, J=7.0 Hz), 4.19-4.31 (2H, m), 4.84 (2H, s), 6.91 (2H, d, J=8.8 Hz), 6.97 (1H, d, J=8.4 Hz), 7.38 (1H, dd, J=8.4, 2.2 Hz), 7.44-7.57 (5H, m), 7.69 (1H, d, J=2.2 Hz), 7.80 (2H, d, J=8.4 Hz), 8.01 (1H, s).

IR (KBr) ν: 3440, 1657, 1605, 1520, 1491, 1317, 1240 cm⁻¹

Anal. for C₃₃H₃₉N₂O₄I·1.0H₂O

Calcd: C, 58.93 ; H, 6.14 ; N, 4.16

Found: C, 58.86 ; H, 6.18 ; N, 4.19.

Reference Example 12

To a solution of 7-(4-ethylphenyl)-N-[4-[N-methyl-N-(tetrahydropyran-4-yl)aminomethyl]phenyl]-2,3-dihydro-1-benzoxepine-4-carboxamide (125 mg) in DMF (5 ml) was added methyl iodide (0.04 ml) at room temperature, and the mixture was stirred for 20 hours. Under reduced pressure, the mixture was concentrated, and to the residue was added ethyl acetate to precipitate solid, which was collected by filtration and recrystallized from acetone-diethylether→ethanol-diethylether) to give dimethyl-[4-N-[7-(4-ethylphenyl)-2,3-dihydro-1-benzoxepin-4-

carbonyl]aminobenzyl]-4-tetrahydropyranylammonium iodide (68 mg) as pale yellow crystals.

m.p. 156-160 °C

¹H-NMR (200MHz, CDCl₃) δ: 1.25 (3H, t, J=7.6 Hz), 1.69-1.93 (2H, m), 2.13-2.28 (2H, m), 2.66 (2H, q, J=7.6 Hz), 2.95 (6H, s), 3.00-3.09 (2H, m), 3.39-3.56 (2H, m), 4.02-4.34 (5H, m), 4.86 (2H, s), 6.99 (1H, d, J=8.4 Hz), 7.18-7.28 (3H, m), 7.39-7.56 (5H, m), 7.69-7.73 (1H, m), 7.79 (2H, d, J=8.8 Hz), 8.78 (1H, s).

IR (KBr) ν: 3429, 1657, 1301, 1520, 1491, 1412, 1319, 1244, 827 cm⁻¹

Anal. for C₃₃H₃₉N₂O₃I·1.0H₂O

Calcd: C, 60.37 ; H, 6.29 ; N, 4.27

Found: C, 60.40 ; H, 6.24 ; N, 4.10.

15 Reference Example 13

To a solution of N-[4-[N-methyl-N-(tetrahydropyran-4-yl)aminomethyl]phenyl]-7-(4-trifluoromethylphenyl)-2,3-dihydro-1-benzoxepine-4-carboxamide (113.6 mg) in DMF (5 ml) was added methyl iodide (0.04 ml) at room temperature, and the mixture was stirred for 24 hours. Under reduced pressure, the mixture was concentrated, and to the residue was added ethyl acetate to precipitate solid, which was collected by filtration and recrystallized from acetone-diethylether→ethanol-diethyl-ether) to give dimethyl-[4-N-[7-(4-trifluoromethylphenyl)-2,3-dihydro-1-benzoxepin-4-carbonyl]aminobenzyl]-4-tetrahydropyranylammonium iodide (99 mg) as pale yellow crystals. m.p. 213 °C (dec.)

¹H-NMR (200MHz, DMSO-d₆) δ: 1.42-1.66 (2H, m), 1.75-1.88 (2H, m), 2.55 (6H, s), 2.62-2.72 (2H, m), 2.94-3.35 (3H, m), 3.68-3.81 (2H, m), 3.96-4.08 (2H, m), 4.13 (2H, s), 6.80 (1H, d, J=8.8 Hz), 7.05 (1H, s), 7.21 (2H, d, J=8.4 Hz), 7.34-7.40 (1H, m), 7.44-7.63 (7H, m), 9.89 (1H, s).

IR (KBr) ν: 3277, 1649, 1510, 1520, 1491, 1325, 1255, 1120, 843 cm⁻¹

Anal. for C₃₂H₃₄N₂O₃F₃I·0.2H₂O

Calcd: C, 56.35 ; H, 5.08 ; N, 4.11

Found: C, 56.21 ; H, 5.16 ; N, 4.11.

Reference Example 14

In 1,2-dichloroethane(400 ml) was suspended p-nitro-
5 benzylamine hydrochloride (30.8 g), 1,4-cyclohexane-dione
monoethyleneketal (25.4 g) and triethylamine (23 ml), and
to the suspension was added sodium triacetoxy boron hydride
(50.9 g) under ice-cooling. Under nitrogen atmosphere, the
mixture was stirred at room temperature for 2.5 hours. Under
10 ice-cooling, 37% formalin (14.6 ml) and sodium triacetoxy
boron hydride (50.9 g) were added to the mixture. Under
nitrogen atmosphere, the mixture was stirred at room
temperature overnight. The mixture was neutralized with
sodium hydrogen carbonate and extracted with 1,2-
15 dichloroethane. The organic layer was washed with sodium
chloride solution and dried with anhydrous magnesium sulfate.
Under reduced pressure, the solvent was evaporated to give
yellow solid (47.5 g), 44 g of which was dissolved in (660
ml). To the mixture was added reduced iron (32 g) little
20 by little, and the mixture was stirred at room temperature
overnight. The solvent was evaporated, and to the residue
was added ethyl acetate. The precipitate was filtered off,
and the filtrate was made alkaline with potassium carbonate
and extracted with ethyl acetate. The organic layer was
25 washed with water and saturated sodium chloride solution
and dried with anhydrous magnesium sulfate. Under reduced
pressure, the solvent was evaporated, and the residue was
purified with silica gel column chromatography (ethyl
acetate/triethylamine/methanol) to give 4-((N-(4,4-
30 ethylenedioxycyclohexyl)-N-methyl)aminomethyl)aniline
(34.1 g) as brown oil.

$^1\text{H-NMR}(\text{CDCl}_3)$ δ : 1.36-1.93 (8H, m), 2.17 (3H, s), 2.43-2.57
(1H, m), 3.46 (2H, s), 3.60 (2H, br), 3.94 (4H, s), 6.64
(2H, d, $J=8.4\text{Hz}$), 7.09 (2H, d, $J=8.4\text{Hz}$).

35 IR(neat) ν : 2946, 1615 cm^{-1} .

Reference Example 15

In dichloromethane (400 ml) was suspended 7-(4-methylphenyl)-2,3-dihydro-1-benzoxepine-4-carboxylic acid (17.0 g), and to the suspension were added oxalyl chloride (10.3 ml) and dimethylformamide (catalytic amount) under ice-cooling. The mixture was stirred at room temperature for 2 hours, and the solvent was evaporated. The residue was dissolved in tetrahydrofuran (300 ml), and the mixture was dropwise added to a solution of 4-((N-(4,4-ethylenedioxcyclohexyl)-N-methyl)aminomethyl)-aniline (16.75 g) and triethylamine (25 ml) in tetrahydrofuran (200 ml), under ice-cooling. Under nitrogen atmosphere, the mixture was stirred at room temperature overnight, and the solvent was evaporated. To the residue was added water, and the mixture was extracted with ethyl acetate. The organic layer was washed with water and saturated sodium chloride solution and dried with anhydrous magnesium sulfate. Under reduced pressure, the solvent was evaporated to give crude crystals, which were recrystallized from ethyl acetate to give N-(4-((N-(4,4-ethylenedioxcyclohexyl)-N-methyl)aminomethyl)phenyl)-7-(4-methylphenyl)-2,3-dihydro-1-benzoxepine-4-carboxamide (17.1 g) as colorless crystals.

mp 192-193°C.

$^1\text{H-NMR}(\text{CDCl}_3) \delta$: 1.48-1.86 (8H, m), 2.20 (3H, s), 2.39 (3H, s), 2.45-2.60 (1H, m), 3.08 (2H, t, $J=4.5\text{Hz}$), 3.56 (2H, s), 3.95 (4H, s), 4.36 (2H, t, $J=4.5\text{Hz}$), 7.06 (1H, d, $J=8.4\text{Hz}$), 7.23-7.33 (4H, m), 7.44-7.56 (7H, m).

IR(KBr) ν : 2948, 1651 cm^{-1} .

Anal. for $\text{C}_{34}\text{H}_{38}\text{N}_2\text{O}_4$:

Calcd: C, 75.81; H, 7.11; N, 5.20.

Found: C, 75.51; H, 6.99; N, 5.29.

Reference Example 16

In acetic acid (100 ml) and 1N hydrochloric acid (200 ml) was dissolved N-(4-((N-(4,4-ethylenedioxcyclohexyl)-N-methyl)aminomethyl)phenyl)-7-(4-methylphenyl)-2,3-dihydro-1-benzoxepine-4-carboxamide (17.1 g), and the

mixture was stirred at 100°C for 1.5 hours and concentrated. The residue was neutralized with 1N sodium hydroxide and extracted with ethyl acetate. The organic layer was washed with water and saturated sodium chloride solution and dried with anhydrous magnesium sulfate. Under reduced pressure, the solvent was evaporated to give crude crystals, which were recrystallized from ethyl acetate-methanol to give N-(4-((N-(4-oxocyclohexyl)-N-methyl)aminomethyl)-phenyl)-7-(4-methylphenyl)-2,3-dihydro-1-benzoxepine-4-carboxamide (12 g) as colorless crystals.
mp 149-150°C.

¹H-NMR(CDCl₃) δ: 1.78-2.13 (4H, m), 2.23 (3H, s), 2.25-2.35 (2H, m), 2.39 (3H, s), 2.45-2.57 (2H, m), 2.84-2.94 (1H, m), 3.08 (2H, t, J=4.4Hz), 3.59 (2H, s), 4.35 (2H, t, J=4.4Hz), 7.06 (1H, d, J=8.0Hz), 7.22-7.34 (4H, m), 7.43-7.57 (6H, m), 7.65 (1H, s).

IR(KBr) ν: 2946, 1713cm⁻¹.

Anal. for C₃₂H₃₄N₂O₃

Calcd: C, 77.70; H, 6.93; N, 5.66.

Found: C, 77.45; H, 6.78; N, 5.65.

Reference Example 17

To a mixture of methyl 2-bromo-6,7-dihydro-5H-benzocycloheptene-8-carboxylate (0.5 g), 4-(1-pyrrolidinyl)phenyl borate(0.37 g), 1M potassium carbonate (6 ml) and ethanol (6 ml) was added toluene (50 ml), and the mixture was stirred under argon atmosphere at room temperature for 30 minutes. To the mixture was added tetrakis(triphenylphosphine)palladium (0.08 g), and the mixture was refluxed for 6 hours and extracted with ethyl acetate. The organic layer was washed with water and saturated sodium chloride solution and dried with anhydrous magnesium sulfate. Under reduced pressure, the solvent was evaporated, and the residue was purified with silica gel column (ethyl acetate/hexane) to give colorless crystals (0.48 g), which were dissolved in 1N sodium hydroxide (15 ml), methanol (50 ml) and tetrahydrofuran (50 ml). The

mixture was stirred at room temperature overnight, concentrated and neutralized with hydrochloric acid to precipitate 2-(4-(1-pyrrolidinyl)phenyl)-6,7-dihydro-5H-benzocycloheptene-8-carboxylic acid (0.46 g) as pale yellow crystals.

mp 242-243°C(dec.).

¹H-NMR(DMSO-d₆) δ: 1.93-2.00 (6H, m), 2.56 (2H, t, J=5.8Hz), 2.76-2.82 (2H, m), 3.23-3.35 (4H, m), 6.60 (2H, d, J=8.8Hz), 7.20 (1H, d, J=8.2Hz), 7.44 (1H, dd, J=1.0, 8.2Hz), 7.53 (2H, d, J=8.8Hz), 7.56 (1H, d, J=1.0Hz), 7.69 (1H, s).

Anal. for C₂₂H₂₃NO₂·0.1H₂O:

Calcd: C, 78.82; H, 6.98; N, 4.18.

Found: C, 78.92; H, 6.95; N, 4.15.

Reference Example 18

To a solution of 2-(4-(1-pyrrolidinyl)phenyl)-6,7-dihydro-5H-benzocycloheptene-8-carboxylic acid (0.45 g), 4-(N-methyl-N-(tetrahydropyran-4-yl)aminomethyl)aniline (0.33 g) and 1-hydroxybenzotriazole (0.18 g) in dimethylformamide (20 ml) was added 1-ethyl-3-(3-dimethylamino-propyl)carbodiimide hydrochloride (0.39 g) under ice-cooling. Under nitrogen atmosphere, the reaction mixture was cooled to room temperature, and to the mixture were added 4-dimethylaminopyridine (catalytic amount) and triethylamine (0.56 ml). The mixture was stirred overnight, poured into water and extracted with ethyl acetate. The organic layer was washed with water and saturated sodium chloride solution and dried with anhydrous magnesium sulfate.

Under reduced pressure, the solvent was evaporated, and the residue was purified with silica gel column (ethyl acetate/methanol/triethylamine) to give crude crystals, which were recrystallized from ethyl acetate-hexane to give 2-(4-(1-pyrrolidinyl)phenyl)-N-(4-(N-tetrahydropyran-4-yl-N-methyl)aminomethyl)phenyl)-6,7-dihydro-5H-benzocycloheptene-8-carboxamide (0.28 g) as colorless crystals.

mp 124-125°C.

$^1\text{H-NMR}(\text{CDCl}_3) \delta$: 1.66-1.77 (4H, m), 1.99-2.06 (4H, m), 2.11-2.18 (2H, m), 2.21 (3H, s), 2.55-2.75 (3H, m), 2.84-2.90 (2H, m), 3.30-3.44 (6H, m), 3.58 (2H, s), 4.00-4.14 (2H, m), 6.64 (2H, d, $J=9.0\text{Hz}$), 7.19 (1H, d, $J=8.0\text{Hz}$), 7.31 (2H, d, $J=8.5\text{Hz}$), 7.39-7.51 (4H, m), 7.57 (2H, d, $J=8.5\text{Hz}$), 7.64 (1H, s).

IR(KBr) ν : 2946, 2843, 1651, 1611 cm^{-1} .

Anal. for $\text{C}_{35}\text{H}_{41}\text{N}_3\text{O}_2 \cdot 0.2\text{H}_2\text{O}$

Calcd: C, 77.95; H, 7.74; N, 7.79.

10 Found: C, 77.76; H, 7.59; N, 7.79.

Reference Example 19

In 1,2-dichloroethane (50 ml) were dissolved p-nitrobenzaldehyde (5 g) and 3-amino-1-propanol (2.5 g), and to the mixture was added sodium triacetoxy boron hydride (9.8 g) under ice-cooling. Under nitrogen atmosphere, the mixture was stirred at room temperature for 5 hours. Under ice-cooling, to the mixture was added 37% formalin (3 ml) and sodium triacetoxy boron hydride (9.8 g). Under nitrogen atmosphere, the mixture was stirred at room temperature overnight. To the mixture was added water, and the mixture was concentrated, neutralized with aqueous sodium hydroxide and extracted with ethyl acetate. The organic layer was washed with water and sodium chloride solution and dried with anhydrous magnesium sulfate. Under reduced pressure, the solvent was evaporated, and the residue was purified with silica gel column (ethyl acetate/methanol/triethylamine) to give yellow oil (5.0 g), 2.5g of which was dissolved in ethanol (50 ml) and catalytic hydrogenation was carried out with 5% palladium on carbon (0.2 g) for 1.5 hours. The catalyst was filtered off, and the solvent was evaporated. The residue was purified with silica gel column (ethyl acetate/methanol/triethylamine) to give 4-((N-3-hydroxypropyl-N-methyl)aminomethyl)-aniline (1.5 g) as pale yellow oil.

35 $^1\text{H-NMR}(\text{CDCl}_3) \delta$: 1.67-1.78 (2H, m), 2.21 (3H, s), 2.62 (2H, t, $J=5.5\text{Hz}$), 3.41 (2H, s), 3.65 (2H, br), 3.77 (2H, t,

J=5.1Hz), 6.65 (2H, d, J=8.4Hz), 7.07 (2H, d, J=8.4Hz).
IR(neat) ν : 3347, 2948, 2799, 1615cm⁻¹.

Reference Example 20

In dichloromethane (5 ml) was suspended 2-(4-methyl-
5 phenyl)-6,7-dihydro-5H-benzocycloheptene-8-carboxylic
acid (0.3 g), and to the suspension were added oxalyl
chloride (0.28 ml) and dimethylformamide (catalytic
amount) under ice-cooling. The mixture was stirred at room
temperature for 1.5 hours, and the solvent was evaporated.
10 The residue was dissolved in tetrahydrofuran (15 ml), and
the mixture was dropwise added to a solution of 4-((N-
3-hydroxypropyl-N-methyl)aminomethyl)aniline (0.23 g) and
triethylamine (0.45 ml) in tetrahydrofuran (15 ml) under
ice-cooling. Under nitrogen atmosphere, the mixture was
15 stirred at room temperature overnight, and the solvent was
evaporated. To the residue was added water, and the mixture
was extracted with ethyl acetate. The organic layer was
washed with water and saturated sodium chloride solution
and dried with anhydrous magnesium sulfate. Under reduced
20 pressure, the solvent was evaporated, and the residue was
purified with silica gel column (ethyl acetate/methanol/
triethylamine) to give crude crystals, which were
recrystallized from ethyl acetate-hexane to give N-(4-
((N-3-hydroxypropyl-N-methyl)aminomethyl)phenyl)-2-(4-
25 methylphenyl)-6,7-dihydro-5H-benzocycloheptene-8-
carboxamide (0.32 g) as colorless crystals.
mp 139-140°C.

¹H-NMR(CDCl₃) δ : 1.72-1.81 (2H, m), 2.13-2.19 (2H, m), 2.25
(3H, s), 2.40 (3H, s), 2.63-2.75 (4H, m), 2.86-2.92 (2H,
30 m), 3.53 (2H, s), 3.79 (2H, t, J=5.4Hz), 7.21-7.32 (3H, m),
7.42-7.52 (6H, m), 7.58 (2H, d, J=8.4Hz), 7.66 (1H, s).
IR(KBr) ν : 2936, 1651cm⁻¹.

Anal. for C₃₀H₃₄N₂O₂·0.5H₂O:

Calcd: C, 77.72; H, 7.61; N, 6.04.

35 Found: C, 77.94; H, 7.62; N, 6.15.

Reference Examl 21

- In dichloromethane(12 ml) was suspended 7-(4-methyl-phenyl)-2,3-dihydro-1-benzoxepine-4-carboxylic acid (0.4 g), and to the suspension were added oxalyl chloride (0.37 ml) and dimethylformamide (catalytic amount) under ice-cooling. The mixture was stirred at room temperature for 2 hours, and the solvent was evaporated. The residue was dissolved in tetrahydrofuran (15 ml), and the mixture was dropwise added to a solution of 4-((N-3-hydroxy-propyl-N-methyl)aminomethyl)aniline (0.33 g) and tri-ethylamine (0.6 ml) in tetrahydrofuran(15 ml) under ice-cooling. Under nitrogen atmosphere, the mixture was stirred at room temperature overnight, and the solvent was evaporated. To the residue was added water, and the mixture was extracted with ethyl acetate. The organic layer was washed with water and saturated sodium chloride solution and dried with anhydrous magnesium sulfate. Under reduced pressure, the solvent was evaporated, and the residue was purified with silica gel column (ethyl acetate/methanol/triethylamine) to give crude crystals, which were recrystallized from ethyl acetate-hexane to give N-(4-((N-3-hydroxypropyl-N-methyl)aminomethyl)phenyl)-7-(4-methylphenyl)-2,3-dihydro-1-benzoxepine-4-carboxamide (0.39 g) as colorless crystals.
- mp 119-120°C.
- ¹H-NMR(CDCl₃) δ: 1.68-1.80 (2H, m), 2.24 (3H, s), 2.39 (3H, s), 2.65 (2H, t, J=5.8Hz), 3.07 (2H, t, J=4.6Hz), 3.52 (2H, s), 3.77 (2H, t, J=5.2Hz), 4.35 (2H, t, J=4.6Hz), 7.05 (1H, d, J=8.4Hz), 7.22-7.31 (3H, m), 7.43-7.52 (5H, m), 7.57 (2H, d, J=8.4Hz), 7.78 (1H, s).
- IR(KBr) ν: 3287, 2948, 1649cm⁻¹.
- Anal. for C₂₉H₃₂N₂O₃·0.2H₂O:
Calcd: C, 75.69; H, 7.10; N, 6.09.
Found: C, 75.58; H, 6.93; N, 6.08.
- Reference Example 22
- In dichloromethane (10 ml) was suspended 7-(4-methyl-phenyl)-2,3-dihydro-1-benzothiepine-4-carboxylic acid

(0.3 g), and to the suspension were added oxalyl chloride (0.27 ml) and dimethylformamide (catalytic amount) under ice-cooling. The mixture was stirred at room temperature for 2 hours, and the solvent was evaporated. The residue
5 was dissolved in tetrahydrofuran (15 ml), and the mixture was dropwise added to a solution of 4-(N-methyl-N-(tetrahydropyran-4-yl)aminomethyl)aniline (0.25 g) and triethylamine (0.42 ml) in tetrahydrofuran (15 ml) under ice-cooling. Under nitrogen atmosphere, the mixture was
10 stirred at room temperature overnight, and the solvent was evaporated. To the residue was added water, and the mixture was extracted with ethyl acetate. The organic layer was washed with water and saturated sodium chloride solution and dried with anhydrous magnesium sulfate. Under reduced
15 pressure, the solvent was evaporated to give crude crystals, which were recrystallized from ethyl acetate-hexane to give 7-(4-methylphenyl)-N-(4-((N-tetrahydropyran-4-yl-N-methyl)aminomethyl)phenyl)-2,3-dihydro-1-benzothiepine-4-carboxamide (0.45 g) as colorless crystals.

20 mp 177-178°C.

$^1\text{H-NMR}(\text{CDCl}_3) \delta$: 1.63-1.77 (4H, m), 2.21 (3H, s), 2.40 (3H, s), 2.57-2.70 (1H, m), 3.08 (2H, t, $J=5.8\text{Hz}$), 3.26-3.44 (4H, m), 3.57 (2H, s), 4.01-4.11 (2H, m), 7.24-7.34 (3H, m), 7.40-7.57 (8H, m), 7.70 (1H, s).

25 IR(KBr) ν : 2949, 1651 cm^{-1} .

Anal. for $\text{C}_{31}\text{H}_{34}\text{N}_2\text{O}_2\text{S} \cdot 0.3\text{H}_2\text{O}$:

Calcd: C, 73.86; H, 6.92; N, 5.56.

Found: C, 73.93; H, 6.73; N, 5.82.

Reference Example 23

30 In dichloromethane (6 ml) was suspended 2-(4-methylphenyl)-6,7-dihydro-5H-benzocycloheptene-8-carboxylic acid (0.25 g), and to the suspension were added oxalyl chloride (0.24 ml) and dimethylformamide (catalytic amount) under ice-cooling. The mixture was stirred at room
35 temperature for 1.5 hours, and the solvent was evaporated. The residue was dissolved in tetrahydrofuran (15 ml, and

th mixture was dropwis added to a solution of 4-((N-methyl-N-(pentan-3-yl))aminomethyl)aniline (0.2 g) and triethylamine (0.38 ml) in tetrahydrofuran (15 ml) under ice-cooling. Under nitrogen atmosphere, the mixture was stirred at room temperature for 5 hours, and the solvent was evaporated. To the residue was added water, and the mixture was extracted with ethyl acetate. The organic layer was washed with water and saturated sodium chloride solution and dried with anhydrous magnesium sulfate. Under reduced pressure, the solvent was evaporated to give crude crystals, which were recrystallized from ethyl acetate-hexane to give N-(4-((N-methyl-N-(pentan-3-yl))aminomethyl)phenyl)-2-(4-methylphenyl)-6,7-dihydro-5H-benzocycloheptene-8-carboxamide (0.23 g) as colorless crystals.

mp 112-113°C.

$^1\text{H-NMR}(\text{CDCl}_3) \delta$: 0.94 (6H, t, $J=7.3\text{Hz}$), 1.26-1.54 (4H, m), 2.14 (3H, s), 2.14-2.32 (3H, m), 2.40 (3H, s), 2.72 (2H, t, $J=6.4\text{Hz}$), 2.86-2.91 (2H, m), 3.55 (2H, s), 7.21-7.27 (3H, m), 7.31-7.56 (8H, m), 7.62 (1H, s).

IR(KBr) ν : 2930, 1651 cm^{-1} .

Anal. for $\text{C}_{32}\text{H}_{38}\text{N}_2\text{O}$:

Calcd: C, 82.36; H, 8.21; N, 6.00.

Found: C, 82.30; H, 8.05; N, 5.90.

Reference Example 24

To a mixture of 3-(4-methylphenyl)-6,7,8,9-tetrahydro-5H-benzocycloheptan-5-one (0.5 g), potassium carbonate (1.65 g) and 18-crown-6 (1.05 g) was added dimethylsulfoxide (10 ml). Under carbon dioxide atmosphere, the mixture was stirred at room temperature for 20 hours, poured into water, acidified with hydrochloric acid and extracted with ethyl acetate. The organic layer was washed with water and subjected to back extraction with sodium hydroxide and water. The aqueous layer was collected, acidified with hydrochloric acid and extracted with ethyl acetate. The organic layer was washed with water and saturated sodium chloride solution and dried with anhydrous

magnesium sulfate. The solvent was evaporated to precipitate colorless crystals (0.42 g), which were filtered with hexane and dissolved in ethanol (40 ml). To the mixture was added sodium boron hydride (0.54 g), and the mixture
5 was stirred at room temperature for 1 hour. To the mixture was added water, and the mixture was concentrated, was acidified with hydrochloric acid and extracted with ethyl acetate. The organic layer was washed with water and saturated sodium chloride solution and dried with anhydrous
10 magnesium sulfate. The solvent was evaporated to give colorless crystals (0.41 g), which were dissolved in 80% formic acid (40 ml). The mixture was stirred at 100°C for 2.5 hours and concentrated. To the residue was added water, and the mixture was extracted with ethyl acetate. The
15 organic layer was washed with water and saturated sodium chloride solution and dried with anhydrous magnesium sulfate. The solvent was evaporated, and the residue was purified with silica gel column (ethyl acetate/hexane) to give
20 2-(4-methylphenyl)-6,7-dihydro-5H-benzocycloheptene-8-carboxylic acid (0.14 g) as colorless crystals.

¹H-NMR(CDCl₃) δ: 2.04-2.18 (2H, m), 2.40 (3H, s), 2.70 (2H, t, J=6.8Hz), 2.86-2.91 (2H, m), 7.21-7.28 (3H, m), 7.44-7.56 (4H, m), 7.91 (1H, s).

25 Reference Example 25

In dimethylsulfoxide (15 ml) were dissolved 3-(4-methylphenyl)-6,7,8,9-tetrahydro-5H-benzocycloheptan-5-one (0.5 g) and 18-crown-6 (1.05 g). Under ice-cooling, potassium t-butoxide (1.65 g) was added to the solution.
30 Under carbon dioxide atmosphere, the mixture was stirred at room temperature for 3 hours, poured into water, acidified with hydrochloric acid and extracted with ethyl acetate. The organic layer was washed with water and subjected to back extraction with sodium hydroxide and water. The
35 aqueous layer was collected, acidified with hydrochloric acid and extracted with ethyl acetate. The organic layer

was washed with water and saturated sodium chloride solution and dried with anhydrous magnesium sulfate. The solvent was evaporated to precipitate colorless crystals (0.47 g), which were filtered with hexane and dissolved in ethanol (40 ml).

5 To the mixture was added sodium boron hydride (0.58 g), and the mixture was stirred at room temperature for 1 hour. To the mixture was added water, and the mixture was concentrated, acidified with hydrochloric acid and extracted with ethyl acetate. The organic layer was washed with water and

10 saturated sodium chloride solution and dried with anhydrous magnesium sulfate. The solvent was evaporated to precipitate colorless crystals (0.46 g), which were filtered with hexane. To the crystals was added 80% formic acid (10ml), and the mixture was refluxed for 1.5 hours. To the

15 mixture was added water, and the mixture was extracted with ethyl acetate. The organic layer was washed with water and subjected to back extraction with sodium hydroxide and water. The aqueous layer was collected, acidified with

20 hydrochloric acid and extracted with ethyl acetate. The organic layer was washed with water and saturated sodium chloride solution and dried with anhydrous magnesium sulfate. The solvent was evaporated to precipitate 2-(4-methylphenyl)-6,7-dihydro-5H-benzocycloheptene-8-carboxylic acid (0.22 g) as colorless

25 crystals.

¹H-NMR(CDCl₃) δ: 2.04-2.16 (2H, m), 2.40 (3H, s), 2.69 (2H, t, J=6.7Hz), 2.86-2.91 (2H, m), 7.21-7.278 (3H, m), 7.44-7.56 (4H, m), 7.89 (1H, s).

Reference Example 26

30 In dimethylformamide (100 ml) was dissolved 7-(4-methylphenyl)-N-(4-((N-(4-oxocyclohexyl)-N-methyl)-aminomethyl)-phenyl)-2,3-dihydro-1-benzoxepin-4-carboxamide (7.5 g), and to the mixture was added methyl iodide (4.7 ml). Under nitrogen atmosphere, the mixture was

35 stirred at room temperature overnight. The solvent was evaporated, and to the residue was added acetone to give

dimethyl-(N-(7-(4-methylphenyl)-2,3-dihydro-1-benzoxepin-4-carbonyl)-4-aminobenzyl)-N-(4-oxocyclohexyl)ammonium iodide (8.9 g) as colorless crystals.

¹H-NMR(DMSO-d₆) δ: 2.09-2.24 (2H, m), 2.34 (3H, s), 2.41-2.61 (6H, m), 2.97 (6H, s), 2.97-3.00 (2H, m), 3.79-3.90 (1H, m), 4.31 (2H, t, J=4.4Hz), 4.56 (2H, s), 7.07 (1H, d, J=8.4Hz), 7.27 (2H, d, J=8.2Hz), 7.37 (1H, s), 7.55-7.60 (5H, m), 7.75 (1H, d, J=2.2Hz), 7.88 (2H, d, J=8.8Hz), 10.20 (1H, s).

10 Reference Example 27

In dimethylformamide (5 ml) was dissolved in 2-(4-(1-pyrrolidinyl)phenyl)-N-(4-((N-tetrahydropyran-4-yl)-N-methyl)aminomethyl)phenyl)-6,7-dihydro-5H-benzocycloheptene-8-carboxamide (0.15 g), and to the mixture was added methyl iodide (0.02 ml). Under nitrogen atmosphere, the mixture was stirred at room temperature overnight. To the mixture was added ethyl acetate, and crude crystal was filtered. The crude crystal was recrystallized from ethanol-ethyl acetate to give dimethyl-(N-(2-(4-(1-pyrrolidinyl)phenyl)-6,7-dihydro-5H-benzocycloheptene-8-carbonyl)-4-aminobenzyl)-4-tetrahydropyranylammonium iodide (0.05 g) as pale brown powder.

¹H-NMR(DMSO-d₆) δ: 1.80-2.20 (10H, m), 2.63 (2H, t, J=5.6Hz), 2.81-2.84 (2H, m), 2.88 (6H, s), 3.24-3.44 (6H, m), 3.54-3.65 (1H, m), 4.02-4.11 (2H, m), 4.46 (2H, s), 6.62 (2H, d, J=9.0Hz), 7.25 (1H, d, J=7.8Hz), 7.36-7.60 (7H, m), 7.88 (2H, d, J=8.4Hz), 10.22 (1H, s).

IR(KBr) ν: 2967, 1663, 1609cm⁻¹.

Anal. for C₃₆H₄₄IN₃O₂·H₂O:

30 Calcd: C, 62.15; H, 6.66; N, 6.04.

Found: C, 61.89; H, 6.30; N, 5.97.

Reference Example 28

In dimethylformamide (5 ml) was dissolved N-(4-((N-3-hydroxypropyl)-N-methyl)aminomethyl)phenyl)-2-(4-methylphenyl)-6,7-dihydro-5H-benzocycloheptene-8-carboxamide (0.2 g), and to the mixture was added methyl iodide (0.04

ml). Under nitrogen atmosphere, the mixture was stirred at room temperature overnight. The solvent was evaporated, and to the residue was added ethyl acetate to give crude crystals, which were filtered and recrystallized from ethanol-ethyl acetate to give N-(3-hydroxypropyl)-N,N-dimethyl-(N-(2-(4-methylphenyl)-6,7-dihydro-5H-benzocycloheptene-8-carbonyl)-4-aminobenzyl)ammonium iodide (0.05 g) as colorless crystals.

mp 210-213°C.

¹H-NMR(CDCl₃+CD₃OD) δ: 2.00-2.20 (4H, m), 2.40 (3H, s), 2.71 (2H, t, J=6.6Hz), 2.87-2.92 (2H, m), 3.10 (6H, s), 3.54-3.65 (2H, m), 3.73 (2H, t, J=5.3Hz), 4.63 (2H, s), 7.22-7.27 (3H, m), 7.43-7.58 (7H, m), 7.80 (2H, d, J=8.4Hz), 9.21 (1H, s). IR(KBr) ν: 3337, 2934, 1653cm⁻¹.

Anal. for C₃₁H₃₇IN₂O₂·0.5H₂O:

Calcd: C, 61.49; H, 6.33; N, 4.63.

Found: C, 61.55; H, 6.22; N, 4.74.

Reference Example 29

In dimethylformamide (5 ml) was dissolved N-(4-((N-3-hydroxypropyl-N-methyl)aminomethyl)phenyl)-7-(4-methylphenyl)-2,3-dihydro-1-benzoxepine-4-carboxamide (0.14 g), and to the mixture was added methyl iodide (0.04 ml). Under nitrogen atmosphere, the mixture was stirred at room temperature overnight. The solvent was evaporated, and to the residue was added ethyl acetate to give crude crystals, which were filtered and recrystallized from ethanol-ethyl acetate to give dimethyl-3-hydroxypropyl-(N-(7-(4-methylphenyl)-2,3-dihydro-1-benzoxepin-4-carbonyl)-4-aminobenzyl)ammonium iodide (0.15 g) as colorless crystals. mp 216-219°C.

¹H-NMR(CDCl₃+CD₃OD) δ: 2.00-2.20 (2H, m), 2.40 (3H, s), 3.06-3.10 (2H, m), 3.10 (6H, s), 3.51-3.61 (2H, m), 3.73 (2H, t, J=5.4Hz), 4.37 (2H, t, J=4.6Hz), 4.61 (2H, s), 7.07 (1H, d, J=8.4Hz), 7.25 (2H, d, J=8.2Hz), 7.46-7.59 (7H, m), 7.81 (2H, d, J=8.2Hz), 9.54 (1H, s). IR(KBr) ν: 3306, 1651cm⁻¹.

Anal. for $C_{30}H_{35}IN_2O_3 \cdot 0.5H_2O$:

Calcd: C, 59.31; H, 5.97; N, 4.61.

Found: C, 59.36; H, 5.95; N, 4.75.

Reference Example 30

- 5 In dimethylformamide (5 ml) was dissolved 7-(4-methylphenyl)-N-(4-((N-tetrahydropyran-4-yl-N-methyl)-aminomethyl)-phenyl)-2,3-dihydro-1-benzothiepine-4-carboxamide (0.19 g), and to the mixture was added methyl iodide (0.03 ml). Under nitrogen atmosphere, the mixture
10 was stirred at room temperature overnight. The solvent was evaporated, and to the residue was added ethyl acetate to give crude crystals, which were filtered and recrystallized from ethanol-hexane to give dimethyl-(N-(7-(4-methylphenyl)-2,3-dihydro-1-benzothiepine-4-carbonyl)-4-aminobenzyl)-N-(4-tetrahydropyranyl)ammonium iodide (0.2
15 g) as colorless crystals.
mp 220-222°C(dec.).

- 1H -NMR(DMSO- d_6) δ : 1.78-1.95 (2H, m), 2.05-2.20 (2H, m), 2.35 (3H, s), 2.88 (6H, s), 2.95-3.05 (2H, m), 3.21-3.32 (4H, m), 3.50-3.65 (1H, m), 4.05-4.15 (2H, m), 4.46 (2H, s), 7.29
20 (2H, d, J=8.0Hz), 7.46-7.63 (7H, m), 7.81-7.90 (3H, m), 10.34 (1H, s).

IR(KBr) ν : 2924, 1657 cm^{-1} .

Reference Example 31

- 25 In dimethylformamide (5 ml) was dissolved N-(4-((N-methyl-N-(pentan-3-yl))aminomethyl)phenyl)-2-(4-methylphenyl)-6,7-dihydro-5H-benzocycloheptene-8-carboxamide (0.17 g), and to the mixture was added methyl iodide (0.08 ml). Under nitrogen atmosphere, the mixture
30 was stirred at 45°C overnight. The solvent was evaporated, and to the residue was added ethyl acetate to give crude crystals, which were filtered and recrystallized from ethanol-ethyl acetate to give dimethyl-(N-(2-(4-methylphenyl)-6,7-dihydro-5H-benzocycloheptene-8-carbonyl)-4-aminobenzyl)-N-(pentan-3-yl)ammonium iodide (0.15 g) as
35 colorless crystals.

mp 190-194°C(dec.).

¹H-NMR(CDCl₃) δ: 1.15 (6H, t, J=7.4Hz), 1.67-1.82 (2H, m),
2.05-2.25 (4H, m), 2.39 (3H, s), 2.73 (2H, t, J=6.6Hz),
2.80-2.90 (2H, m), 3.11 (6H, s), 3.40-3.51 (1H, m), 4.91
5 (2H, s), 7.18-7.26 (3H, m), 7.44 (1H, dd, J=1.8, 8.4Hz),
7.49 (2H, d, J=8.4Hz), 7.57-7.62 (4H, m), 7.80 (2H, d,
J=8.4Hz), 8.35 (1H, s).

IR(KBr) ν: 2936, 1659cm⁻¹.

Anal. for C₃₃H₄₁IN₂O·0.5H₂O:

10 Calcd: C, 64.18; H, 6.85; N, 4.54.

Found: C, 63.84; H, 6.73; N, 4.47.

Reference Example 32

In DMF (50 ml) was dissolved N-cyclohexyl-N-methylamine (12.5 g, 0.11 mol), and to the solution were
15 added potassium carbonate (27.6 g, 0.20 mol) and 4-nitrobenzylbromide (21.6 g, 0.10 mol). The mixture was stirred at room temperature for 5 hours. Under reduced pressure, the reaction mixture was concentrated. To the residue was added ethyl acetate, and the mixture was
20 extracted with water. The ethyl acetate layer was washed with saturated sodium chloride solution, dried with MgSO₄ and concentrated under reduced pressure. The residue was purified with silica gel column chromatography (ethyl acetate/hexane) to give N-cyclohexyl-N-methyl-N-(4-
25 nitrobenzyl)amine (24.8 g).

¹H-NMR (200 MHz, CDCl₃) δ: 1.0-1.95 (10H, m), 2.19 (3H, s),
3.66 (2H, s), 7.51 (2H, d, J=8.8Hz), 8.17 (2H, d, J=8.8Hz).

Reference Example 33

To a solution of N-cyclohexyl-N-methyl-N-(4-
30 nitrobenzyl)amine (12.4 g, 50.0 mmol) in methanol (250 ml) were added nickel bromide (1.09 g, 5.0 mmol) and then sodium boron hydride (7.57 g, 200 mmol) at 0°C, and the mixture was stirred at room temperature for 30 minutes. To the mixture were added nickel bromide (0.55 g, 2.5 mmol) and then sodium
35 boron hydride (3.78 g, 100 mmol) at 0°C, and the mixture was stirred at room temperature for 30 minutes. To the reaction

mixture was added water (100 ml), and the mixture was concentrated under reduced pressure. To the residue was added ethyl acetate, and insoluble material was filtered off with Celite. The filtrate was washed with ethyl acetate, and the ethyl acetate layer was dried with MgSO₄ and concentrated under reduced pressure. The residue was washed with hexane to give 4-(N-cyclohexyl-N-methylamino-methyl)aniline (3.99 g, 37%).

¹H-NMR (200 MHz, CDCl₃) δ : 1.0-1.95 (10H, m), 2.17 (3H, s), 2.3-2.55 (1H, m), 3.46 (2H, s), 3.59 (2H, br s), 6.65 (2H, d, J=8.5Hz), 7.10 (2H, d, J=8.5Hz).

Reference Example 34

To a solution of 7-(4-methylphenyl)-2,3-dihydro-1-benzoxepine-4-carboxylic acid (0.28 g), 4-(N-cyclohexyl-N-methylaminomethyl)aniline (0.24 g) and 1-hydroxybenzotriazole (0.15 g) in dimethylformamide (10 ml) was added 1-ethyl-3-(3-dimethylaminopropyl)carbodiimide hydrochloride (0.29 g) under ice-cooling. Under nitrogen atmosphere, the mixture was cooled to room temperature, and to the mixture were added 4-dimethylaminopyridine (3 mg) and triethylamine (0.42 ml). The mixture was stirred for 20 hours, poured into water, and extracted with ethyl acetate. The organic layer was washed with water and saturated sodium chloride solution and dried with anhydrous magnesium sulfate. Under reduced pressure, the solvent was evaporated, and the residue was washed with ethyl acetate and dried to give N-(4-(N-cyclohexyl-N-methylaminomethyl)phenyl)-7-(4-methylphenyl)-2,3-dihydro-1-benzoxepine-4-carboxamide (0.40 g).

¹H-NMR(CDCl₃) δ : 1.0-1.95 (10H, m), 2.20 (3H, s), 2.35-2.55 (1H, m), 2.40 (3H, s), 3.0-3.15 (2H, m), 3.56 (2H, s), 4.3-4.45 (2H, m), 7.06 (1H, d, J=8.4Hz), 7.2-7.6 (11H, m).

Reference Example 35

In dimethylformamide (7 ml) was dissolved N-(4-(N-cyclohexyl-N-methylaminomethyl)phenyl)-7-(4-methylphenyl)-2,3-dihydro-1-benzoxepine-4-carboxamid (0.15 g),

and to the mixture was added methyl iodide (0.06 ml). Under nitrogen atmosphere, the mixture was stirred at room temperature for 20 hours. The solvent was evaporated, and to the residue was added ethyl acetate to give crude crystals, which were filtered and recrystallized from ethanol to give N-cyclohexyl-N,N-dimethyl-N-((7-(4-methylphenyl)-2,3-dihydro-1-benzoxepin-4-carbonyl)-4-aminobenzyl)ammonium iodide (0.15 g).

¹H-NMR(CDCl₃) δ: 1.0-1.8 (6H, m), 1.9-2.05 (2H, m), 2.25-2.45 (2H, m), 2.36 (3H, s), 2.95-3.15 (8H, m), 3.45-3.7 (1H, m), 4.2-4.35 (2H, m), 4.83 (2H, s), 6.99 (1H, d, J=8.4Hz), 7.21 (2H, d, J=7.6Hz), 7.35-7.6 (6H, m), 7.74 (1H, d, J=2.2Hz), 7.85 (2H, d, J=8.6Hz), 8.79 (1H, s).
IR(KBr) ν: 1659, 1609, 1593, 1518, 1493cm⁻¹.

15 Reference Example 36

In dimethylformamide (5 ml) was dissolved N-(4-(N-methyl-N-(tetrahydropyran-4-yl)aminomethyl)phenyl)-7-(4-morpholino-phenyl)-2,3-dihydro-1-benzoxepine-4-carboxamide (0.20 g), and to the mixture was added methyl iodide (0.03 ml). Under nitrogen atmosphere, the mixture was stirred at room temperature for 32 hours. The solvent was evaporated, and the residue was purified with silica gel column chromatography (dichloromethane/methanol). The desired fraction was concentrated, and to the residue was added ethyl acetate. Insoluble material was filtered and recrystallized from ethanol to give dimethyl-N-(7-(4-morpholinophenyl)-2,3-dihydro-1-benzoxepin-4-carbonyl)-4-aminobenzyl-N-(4-tetrahydropyranyl)ammonium iodide (0.18 g).

¹H-NMR(CDCl₃) δ: 1.6-2.0 (2H, m), 2.1-2.3 (2H, m), 2.92 (6H, s), 2.95-3.2 (6H, m), 3.35-3.55 (2H, m), 3.8-3.9 (4H, m), 4.0-4.35 (5H, m), 4.84 (2H, s), 6.85-7.05 (3H, m), 7.35-7.85 (9H, m), 8.92 (1H, s).
IR(KBr) ν: 1659, 1609, 1520, 1495cm⁻¹.

35 Reference Example 37

In tetrahydrofuran(100 ml) was dissolved 1,2-

methylenedioxy-4-bromobenzene (24.0 g), and to the mixture was dropwise added n-butyllithium (1.6M hexane solution, 82 ml) at -55°C or less. The mixture was stirred at -70°C or less for 30 minutes. The resulting solution was dropwise
5 added to a solution of trimethyl borate (18.6 g) in tetrahydrofuran (50 ml) at -60°C or less through cannula, and the mixture was stirred at -70°C or less for 1 hour and then for 2 hours while warming the mixture to room temperature. To the reaction mixture were added 1N
10 hydrochloric acid (130 ml) and diethylether (150 ml), and the organic layer was separated. The organic layer was washed with water and saturated sodium chloride solution and dried with anhydrous magnesium sulfate. Under reduced pressure, the solvent was evaporated. The residue was
15 washed with diisopropylether to give 3,4-methylenedioxyphenyl borate (6.79 g).
¹H-NMR(DMSO-d₆) δ: 5.99 (2H, s), 6.8-6.95 (1H, m), 7.25-7.45 (2H, m).

Reference Example 38

20 To a mixture of methyl 7-bromo-2,3-dihydro-1-benzoxepine-4-carboxylate (0.57 g), 3,4-methylenedioxyphenyl borate (0.47 g) and sodium carbonate (0.42 g) were added water (2 ml) and 1,2-dimethoxyethane (12 ml). Under argon atmosphere, the mixture was stirred at room
25 temperature for 30 minutes, and to the mixture was added tetrakis(triphenylphosphine)palladium (0.16 g). The mixture was stirred at 80°C for 14 hours and extracted with ethyl acetate. The organic layer was washed with water and saturated sodium chloride solution and dried with anhydrous
30 magnesium sulfate. Under reduced pressure, the solvent was evaporated, and the residue was purified with silica gel column (ethyl acetate/hexane) to give methyl 7-(3,4-methylenedioxyphenyl)-2,3-dihydro-1-benzoxepine-4-carboxylate (0.43 g).
35 ¹H-NMR(CDCl₃) δ: 2.95-3.10 (2H, m), 3.83 (3H, s), 4.25-4.35 (2H, m), 6.01 (2H, s), 6.87 (1H, d, J=8.6Hz), 6.95-7.10 (3H,

m), 7.40 (1H, dd, J=8.4, 2.4Hz), 7.47 (1H, d, J=2.2Hz), 7.65 (1H, s).

Reference Example 39

To methyl 7-(3,4-methylenedioxyphenyl)-2,3-dihydro-
5 1-benzoxepine-4-carboxylate (0.40 g) were added methanol (5 ml) and 1N sodium hydroxide (3.7 ml), and the mixture was stirred at room temperature for 20 hours. To the mixture was added 1N hydrochloric acid (3.7 ml), and the mixture was concentrated under reduced pressure. Precipitate was
10 washed with water and diethylether and dried under reduced pressure to give 7-(3,4-methylene-dioxyphenyl)-2,3-dihydro-1-benzoxepine-4-carboxylic acid (0.32 g).

¹H-NMR(DMSO-d₆) δ: 2.80-2.95 (2H, m), 4.15-4.35 (2H, m), 6.05 (2H, s), 6.97 (1H, d, J=8.1Hz), 7.01 (1H, d, J=8.4Hz), 7.16
15 (1H, dd, J=8.1, 1.7Hz), 7.29 (1H, d, J=1.7Hz), 7.53 (2H, dd, J=8.4, 2.3Hz), 7.63 (1H, s), 7.74 (1H, d, J=2.3Hz).

Reference Example 40

To a solution of 7-(3,4-methylenedioxyphenyl)-2,3-dihydro-1-benzoxepine-4-carboxylic acid (0.14 g), 4-(N-methyl-N-(tetrahydropyran-4-yl)aminomethyl)aniline (0.11
20 g) and 1-hydroxy-benzotriazole (0.15 g) in dimethylformamide (10 ml) was added 1-ethyl-3-(3-dimethylaminopropyl)carbodiimide hydrochloride (0.13 g) under ice-cooling. Under nitrogen atmosphere, the reaction
25 mixture was warmed to room temperature. To the mixture were added 4-dimethylaminopyridine (3 mg) and triethylamine (0.19 ml), and the mixture was stirred for 18 hours, poured into water, and extracted with ethyl acetate. The organic layer was washed with water and saturated sodium chloride
30 solution, and dried with anhydrous magnesium sulfate. Under reduced pressure, the solvent was evaporated, and the residue was purified with silica gel column (ethyl acetate) to give 7-(3,4-methylenedioxyphenyl)-4-(N-methyl-N-(tetrahydropyran-4-yl)aminomethyl)phenyl)-2,3-dihydro-
35 1-benzoxepine-4-carboxamide (0.19 g).

¹H-NMR(CDCl₃) δ: 1.55-1.85 (4H, m), 2.21 (3H, s), 2.55-2.80

(1H, m), 3.00-3.15 (2H, m), 3.30-3.45 (2H, m), 3.58 (2H, s), 3.95-4.15 (2H, m), 4.30-4.45 (2H, m), 6.01 (2H, s), 6.88 (1H, d, J=8.6Hz), 6.95-7.10 (3H, m), 7.20-7.65 (7H, m). IR(KBr) ν : 1653, 1597, 1514, 1483cm⁻¹.

5 Reference Example 41

In dimethylformamide (5 ml) was dissolved 7-(3,4-methylenedioxyphenyl)-4-(N-methyl-N-(tetrahydropyran-4-yl)aminomethyl)phenyl)-2,3-dihydro-1-benzoxepine-4-carboxamide (95 mg), and to the mixture was added methyl
10 iodide (0.012 ml). Under nitrogen atmosphere, the mixture was stirred at room temperature for 18 hours. The solvent was evaporated, and to the residue was added ethyl acetate.

Insoluble material was filtered and recrystallized from ethanol to give dimethyl-N-(7-(3,4-methylenedioxy-
15 phenyl)-2,3-dihydro-1-benzoxepin-4-carbonyl)-4-amino-benzyl-N-(4-tetrahydropyranyl)ammonium iodide (101 mg).
¹H-NMR(CDCl₃) δ : 1.7-2.0 (2H, m), 2.15-2.3 (2H, m), 2.85-3.1 (8H, m), 3.4-3.55 (2H, m), 4.0-4.35 (5H, m), 4.85 (2H, s), 5.96 (2H, s), 6.81 (1H, d, J=7.8Hz), 6.9-7.1 (3H, m),
20 7.25-7.7 (5H, m), 7.83 (2H, d, J=8.2 Hz), 8.89 (1H, s).
IR(KBr) ν : 1659, 1609, 1520, 1495cm⁻¹.

Reference Example 42

A mixture of methyl 2-bromo-6,7-dihydro-5H-benzocyclohepten-8-carboxylate (1.0g) in methanol (25ml),
25 THF (25ml) and 1N NaOH (18ml) was refluxed for 1 hour. The solvent was concentrated, acidified using 1N HCl, and extracted with ethyl acetate. The organic layer was washed with water and brine, and dried over MgSO₄. The solvent was evaporated in vacuo to give 2-bromo-6,7-dihydro-5H-
30 benzocyclohepten-8-carboxylic acid (0.95g) as colorless prisms.

mp 213-215°C(dec.).

¹H-NMR(δ ppm, CDCl₃) 2.00-2.12 (2H, m), 2.62-2.68 (2H, m), 2.76-2.82 (2H, m), 7.04 (1H, d, J=8.2Hz), 7.35 (1H, dd, J=2.0,
35 8.2Hz), 7.48 (1H, d, J=2.0Hz), 7.74 (1H, s).
IR(KBr) ν : 2932, 1669cm⁻¹.

Anal. calcd. for $C_{12}H_{11}BrO_2$: C, 53.96; H, 4.15.

Found C, 53.93; H, 4.03.

Reference Example 43

Oxalyl chloride (0.5ml) and DMF (cat.) were added
5 successively to a solution of 2-bromo-6,7-dihydro-5H-
benzocyclohepten-8-carboxylic acid (0.95g) in THF (20ml)
under ice cooling. The mixture was stirred for 1.5 hours
at room temperature, and the solvent was evaporated in vacuo.
A solution of the residue in THF (25ml) was added dropwise
10 to a solution of 4-[[N-methyl-N-(tetrahydropyran-4-
yl)amino]methyl]aniline (0.9g) and triethylamine (1.5ml)
in THF (20ml) under ice cooling. The reaction mixture was
stirred for 1.5 hours at room temperature under nitrogen.
The solvent was evaporated in vacuo, water was added to the
15 residue, and the mixture was extracted with ethyl acetate.
The organic layer was washed with water and brine, and dried
over $MgSO_4$. The solvent was evaporated in vacuo to give
2-bromo-N-[4-[[N-methyl-N-(tetrahydropyran-4-
yl)amino]methyl]phenyl]-6,7-dihydro-5H-
20 benzocyclohepten-8-carboxamide (1.55g) as colorless
prisms.
mp 116-121°C.

1H -NMR (δ ppm, $CDCl_3$) 1.64-1.77 (4H, m), 2.05-2.18 (2H, m),
2.21 (3H, s), 2.59-2.71 (3H, m), 2.76-2.81 (2H, m), 3.37
25 (2H, dt, $J=3.0$, 11.2Hz), 3.57 (2H, s), 4.01-4.07 (2H, m),
7.04 (1H, d, $J=8.0$ Hz), 7.22 (1H, s), 7.29-7.35 (3H, m), 7.42
(1H, d, $J=1.8$ Hz), 7.55 (2H, d, $J=8.4$ Hz), 7.64 (1H, br).
IR (KBr) ν : 3279, 2942, 2845, 1653, 1516 cm^{-1} .

Anal. calcd. for $C_{25}H_{29}BrN_2O_2$: C, 63.97; H, 6.23; N, 5.97.

30 Found C, 63.67; H, 6.10; N, 5.92.

Reference Example 44

A solution of 4-carboxyphenylboronic acid (2.0g) and
sulfuric acid (0.1ml) in methanol (50ml) was refluxed
overnight. The reaction mixture was poured into water, and
35 the mixture was extracted with ethyl acetate. The organic
layer was washed with water and brine, and dried over $MgSO_4$.

The solvent was evaporated in vacuo to give 4-methoxycarbonylphenylboronic acid (1.7g) as colorless prisms.

¹H-NMR(δ ppm, CDCl₃+DMSO-d₆) 3.92 (3H, s), 6.78 (2H, s), 7.93 (2H, d, J=7.8Hz), 8.02 (2H, d, J=7.8Hz).

IR(KBr) ν: 3370, 1709cm⁻¹.

Reference Example 45

A mixture of 2-bromo-N-[4-[[N-methyl-N-(tetrahydropyran-4-yl)amino]methyl]phenyl]-6,7-dihydro-5H-benzocyclohepten-8-carboxamide (0.95g) and 4-methoxycarbonylphenylboronic acid (0.4g) in 1M potassium carbonate (6ml), ethanol (6ml) and toluene (75ml) was stirred for 30 minutes at room temperature under argon. Tetrakis(triphenylphosphine)palladium(0.12g) was added to the mixture, and the mixture was refluxed for 6 hours under argon. The reaction mixture was extracted with ethyl acetate. The organic layer was washed with water and brine, and dried over MgSO₄. The solvent was evaporated in vacuo. The residue was subjected to a silica gel column chromatography (ethyl acetate/methanol/triethylamine) to give 2-(4-methoxycarbonylphenyl)-N-[4-[[N-methyl-N-(tetrahydropyran-4-yl)amino]methyl]phenyl]-6,7-dihydro-5H-benzocyclohepten-8-carboxamide (0.4g) as colorless prisms.

mp 176-183°C.

¹H-NMR(δ ppm, CDCl₃) 1.70-1.77 (4H, m), 2.14-2.21 (2H, m), 2.21 (3H, s), 2.60-2.76 (3H, m), 2.88-2.93 (2H, m), 3.37 (2H, dt, J=2.6, 11.0Hz), 3.57 (2H, s), 3.94 (3H, s), 4.01-4.07 (2H, m), 7.26-7.34 (3H, m), 7.43 (1H, s), 7.49 (1H, dd, J=1.8, 8.0Hz), 7.54-7.57 (3H, m), 7.58-7.68 (3H, m), 8.11 (2H, d, J=8.4Hz).

IR(KBr) ν: 3281, 2948, 2847, 1721cm⁻¹.

Reference Example 46

A mixture of 2-(4-methoxycarbonylphenyl)-N-[4-[[N-methyl-N-(tetrahydropyran-4-yl)amino]methyl]phenyl]-6,7-dihydro-5H-benzocyclohepten-8-carboxamide (0.25g) in

methanol (25ml), THF (25ml) and 1N NaOH (5ml) was refluxed overnight. The reaction mixture was concentrated, and neutralized using 1N HCl. The precipitate was filtered, washed with water and dried to give 2-(4-

- 5 carboxyphenyl)-N-[4-[[N-methyl-N-(tetrahydropyran-4-yl)amino]methyl]phenyl]-6,7-dihydro-5H-benzocyclohepten-8-carboxamide (0.21g) as colorless amorphous.

¹H-NMR (δ ppm, DMSO-d₆) 1.40-1.74 (4H, m), 2.05 (2H, br), 2.10
10 (3H, s), 2.60-2.64 (3H, m), 2.84 (2H, br), 3.20-3.27 (2H, m), 3.51 (2H, s), 3.87-3.93 (2H, m), 7.24 (2H, d, J=8.4Hz), 7.30 (1H, d, J=8.4Hz), 7.40 (1H, s), 7.52-7.70 (6H, m), 7.93 (2H, d, J=8.0Hz), 10.06 (1H, s).

IR(KBr) ν: 2938, 2845, 1647, 1595, 1518, 1395cm⁻¹.

- 15 Reference Example 47

To 2-(4-carboxyphenyl)-N-[4-[[N-methyl-N-(tetrahydropyran-4-yl)amino]methyl]phenyl]-6,7-dihydro-5H-benzocyclohepten-8-carboxamide (0.32g) was added 1N HCl (2ml), and the solvent was evaporated in vacuo, and dried.
20 Diphenyldiazomethane (0.4g) was added to a solution of the residue in DMF (40ml), and the reaction mixture was heated overnight at 50°C under nitrogen. The solvent was evaporated in vacuo, and aq. NaHCO₃ was added to the residue. The mixture was extracted with ethyl acetate. The organic
25 layer was washed with water and brine, and dried over MgSO₄. The solvent was evaporated in vacuo. The residue was subjected to a silica gel column chromatography (ethyl acetate/methanol/triethylamine) to give 2-[4-(diphenylmethoxycarbonyl)phenyl]-N-[4-[[N-methyl-N-(tetrahydropyran-4-yl)amino]methyl]phenyl]-6,7-dihydro-5H-benzocyclohepten-8-carboxamide (0.4g) as colorless
30 prisms.

¹H-NMR (δ ppm, CDCl₃) 1.63-1.76 (4H, m), 2.05-2.20 (2H, m), 2.20 (3H, s), 2.55-2.71 (1H, m), 2.71 (2H, t, J=6.4Hz),
35 2.85-2.90 (2H, m), 3.36 (2H, dt, J=2.6, 11.0Hz), 3.57 (2H, s), 4.00-4.06 (2H, m), 7.14 (1H, s), 7.23-7.66 (20H, m),

7.79 (1H, s), 8.19 (2H, d, J=8.4Hz).

Working Example 1 (Production of Compound 1)

In aqueous methanol was dissolved N,N-dimethyl-N-
5 (4-(((2-(4-methylphenyl)-6,7-dihydro-5H-benzocyclohepten-8-yl)carbonyl)amino)benzyl)-N-(4-tetrahydropyranyl)ammonium iodide (19 g), and the mixture was subjected to ion exchange resin (DOWEX1-x8, 100-200 mesh, Cl⁻ type) column, which was eluted with aqueous methanol.
10 The solvent of the desired fractions was evaporated, and to the residue was added acetone to give crude crystals, which were recrystallized from ethanol to give N,N-dimethyl-N-(4-(((2-(4-methylphenyl)-6,7-dihydro-5H-benzocyclohepten-8-yl)carbonyl)amino)benzyl)-N-(4-
15 tetrahydropyranyl)ammonium chloride (10.1 g) as colorless crystals.
mp 226-232°C(dec.).

¹H-NMR(CDCl₃+CD₃OD) δ: 1.80-2.00 (2H, m), 2.07-2.26 (4H, m),
2.39 (3H, s), 2.72 (2H, t, J=6.6Hz), 2.85-2.91 (2H, m), 3.00
20 (6H, s), 3.54 (2H, t, J=11.3Hz), 4.00-4.21 (3H, m), 4.70 (2H, s), 7.21-7.29 (3H, m), 7.42-7.56 (7H, m), 7.81 (2H, d, J=8.4Hz), 9.06 (1H, s).

IR(KBr) ν: 2934, 1655cm⁻¹.

Anal. for C₃₃H₃₉ClN₂O₂:

25 Calcd: C, 74.62; H, 7.40; N, 5.27; Cl, 6.67.
Found: C, 74.35; H, 7.33; N, 5.20; Cl, 6.80.

Working Example 2 (Production of Compound 1)

To a solution of N-(4-chloromethylphenyl)-2-(4-methylphenyl)-6,7-dihydro-5H-benzocycloheptene-8-
30 carboxamide (9.38 g, 23.3 mmol) in DMF (50 ml) was dropwise added a solution of N,N-dimethyl-N-tetrahydropyran-4-ylamine (4.5 g, 35.0 mmol) in DMF (50 ml). Under nitrogen atmosphere, the mixture was stirred for 23 hours. The solvent was evaporated to give powder, which was washed with
35 acetone and dried. The resulting colorless powder was recrystallized from ethanol to give N,N-dimethyl-N-(4-

(((2-(4-methylphenyl)-6,7-dihydro-5H-benzocyclohepten-8-yl)carbonyl)amino)benzyl)-N-(4-tetrahydropyranyl)ammonium chloride (Compound 1) (10.6 g, 86%) as colorless powder.

5 Working Example 3 (Production of Compound 2)

In aqueous acetonitrile was dissolved N,N-dimethyl-N-(((7-(4-methylphenyl)-2,3-dihydro-1-benzoxepin-4-yl)carbonyl)amino)benzyl)-N-(4-oxocyclohexyl)ammonium iodide (22.8 g), and the mixture was subjected to ion exchange resin (DOWEX-SBR, Cl⁻ type) column, which was eluted with aqueous acetonitrile. The solvent of the desired fractions was evaporated, and the residue was dissolved in water. The mixture was subjected to freeze-drying to give N,N-dimethyl-N-(((7-(4-methylphenyl)-2,3-dihydro-1-benzoxepin-4-yl)carbonyl)amino)benzyl)-N-(4-oxocyclohexyl)ammonium chloride (Compound 2) (16.1 g) as colorless powder.

¹H-NMR(DMSO-d₆) δ: 2.05-2.25 (2H, m), 2.34 (3H, s), 2.41-2.61 (6H, m), 2.97 (6H, s), 2.97-3.00 (2H, m), 3.75-3.90 (1H, m), 4.30 (2H, t, J=4.4Hz), 4.57 (2H, s), 7.06 (1H, d, J=8.4Hz), 7.27 (2H, d, J=7.8Hz), 7.45 (1H, s), 7.53-7.60 (5H, m), 7.78 (1H, d, J=2.2Hz), 7.92 (2H, d, J=8.4Hz), 10.34 (1H, s).

IR(KBr) ν: 3025, 2967, 1717, 1655cm⁻¹.

25 Anal. for C₃₃H₃₇ClN₂O₃·0.5H₂O:

Calcd: C, 71.53; H, 6.91; N, 5.06; Cl, 6.40.

Found: C, 71.21; H, 6.94; N, 4.94; Cl, 6.24.

Working Example 4 (Production of Compound 2)

To a solution of N-(4-chloromethylphenyl)-7-(4-methylphenyl)-2,3-dihydro-1-benzoxepine-4-carboxamide (214 mg, 0.530 mmol) in N,N-dimethylformamide (1 ml) was dropwise added a solution of 4-dimethylaminocyclohexanone (112 mg, 0.795 mmol) in N,N-dimethylformamide (1 ml). Under nitrogen atmosphere, the mixture was stirred for 14 hours.

35 The solvent was evaporated to give crude product, which was washed with ether to give N,N-dimethyl-N-(((7-(4-

methylphenyl)-2,3-dihydro-1-benzoxepin-4-yl)carbonyl)amino)benzyl)-N-(4-oxocyclohexyl)ammonium chloride (Compound 2) (305 mg) as colorless powder.

Working Example 5 (Production of Compound 3)

- 5 To a solution of N-(4-chloromethylphenyl)-7-(4-ethoxyphenyl)-2,3-dihydro-1-benzoxepine-4-carboxamide (2.38 g) in DMF (20 ml) was added N,N-dimethyl-N-tetrahydropyran-4-ylamine (1.42 g) at room temperature, and the mixture was stirred for 14 hours. To the reaction mixture
10 was added ethyl acetate (100 ml) to precipitate crystals, which were collected by filtration. The crystal was washed with ethyl acetate to give crude product as pale yellow crystals, which were recrystallized from ethanol to give
15 as N-(4-(((7-(4-ethoxyphenyl)-2,3-dihydro-1-benzoxepin-4-yl)carbonyl)amino)benzyl)-N,N-dimethyl-N-(4-tetrahydropyranyl)ammonium chloride (Compound 3) (1.29 g) colorless crystals.

m.p. 200-204 °C

- ¹H-NMR (200MHz, DMSO-d₆) δ: 1.35 (3H, t, J=7.0 Hz), 1.75-
20 1.98 (2H, m), 2.06-2.24 (2H, m), 2.88 (6H, s), 2.94-3.05 (2H, m), 3.28-3.43 (2H, m), 3.49-3.69 (1H, m), 3.99-4.13 (2H, m), 4.07 (2H, q, J=7.0 Hz), 4.23-4.35 (2H, m), 4.47 (2H, s), 6.98-7.07 (3H, m), 7.37 (1H, s), 7.50-7.61 (5H, m), 7.72 (1H, d, J=2.2 Hz), 7.87 (2H, d, J=8.4 Hz), 10.22
25 (1H, s).

IR (KBr) ν: 3425, 1647, 1603, 1520, 1489, 1407, 1317, 1294, 1240, 831 cm⁻¹.

Anal. for C₃₃H₃₉N₂O₄Cl

Calcd: C, 70.38 ; H, 6.98 ; N, 4.97 ; Cl, 6.30

- 30 Found: C, 70.49 ; H, 7.08 ; N, 4.94 ; Cl, 6.19.

Working Example 6 (Production of Compound 3)

- In aqueous methanol was dissolved N-(4-(((7-(4-ethoxyphenyl)-2,3-dihydro-1-benzoxepin-4-yl)carbonyl)amino)benzyl)-N,N-dimethyl-N-(4-tetrahydropyranyl)ammonium iodide (26.6 g), and the mixture
35 was subjected to ion exchange resin (DOWEX-SBR, Cl⁻ type)

column, which was eluted with aqueous methanol. The solvent of the desired fractions was evaporated, and to the residue was added acetone to give crude crystals, which were recrystallized from ethanol to give N-(4-(((7-(4-ethoxyphenyl)-2,3-dihydro-1-benzoxepin-4-yl)carbonyl)amino)benzyl)-N,N-dimethyl-N-(4-tetrahydropyranyl)ammonium chloride (Compound 3) (16.6 g) as colorless crystals.

Working Example 7 (Production of Compound 4)

- 10 A solution of 2-[4-(diphenylmethoxycarbonyl)-phenyl]-N-[4-[[N-methyl-N-(tetrahydropyran-4-yl)amino]methyl]phenyl]-6,7-dihydro-5H-benzocyclohepten-8-carboxamide (0.4g) and methyl iodide (0.15ml) in DMF (40ml) was stirred overnight under nitrogen.
- 15 The solvent was evaporated in vacuo. Ethyl acetate was added to the residue, and the precipitate was filtered, and washed with ethyl acetate. Trifluoroacetic acid (12ml) was added to a solution of the precipitate in dichloromethane (25ml), and the reaction mixture was stirred for 2.5 hours
- 20 at room temperature. The solvent was evaporated in vacuo. Ethyl acetate was added to the residue, and the solvent was evaporated in vacuo. A mixture of the residue in 50% acetonitrile was subjected to an ion exchange column (Dowex SBR, Cl⁻ form, eluted with 50% acetonitrile). The eluate
- 25 was evaporated in vacuo to give N-[4-[[[2-(4-carboxyphenyl)-6,7-dihydro-5H-benzocyclohepten-8-yl]carbonyl]amino]benzyl]-N,N-dimethyl-N-(4-tetrahydro-2H-pyranyl)ammonium chloride (Compound 4) (0.23g) as colorless amorphous.
- 30 ¹H-NMR(δ ppm, DMSO-d₆) 1.80-2.20 (6H, m), 2.65 (2H, t, J=7.2Hz), 2.80-2.88 (2H, m), 2.88 (6H, s), 3.24-3.40 (2H, m), 3.53-3.65 (1H, m), 4.05-4.10 (2H, m), 4.47 (2H, s), 7.35-7.41 (2H, m), 7.54 (2H, d, J=8.4Hz), 7.64 (1H, d, J=8.0Hz), 7.80-7.91 (5H, m), 8.03 (2H, d, J=8.4Hz), 10.27
- 35 (1H, s).
- IR(KBr) ν: 3034-2872, 1696, 1659, 1607, 1593, 1518cm⁻¹.

Anal. calcd. for $C_{33}H_{37}ClN_2O_4 \cdot 0.5H_2O$:

C, 69.52; H, 6.72; N, 4.91; Cl, 6.22.

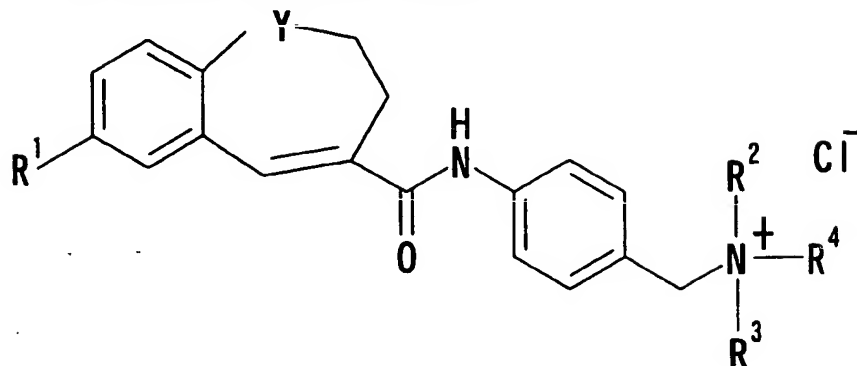
Found C, 69.19; H, 6.80; N, 5.03; Cl, 6.15.

5 Industrial Applicability

The compound of the formula (I) of the present invention has potent CCR5 antagonistic activity and can be advantageously used for the treatment or prevention of infectious disease of various HIV in human (e.g. AIDS).

CLAIMS

1. A compound of the formula:



- 5 wherein R¹ is an optionally substituted phenyl or an optionally substituted thienyl; Y is -CH₂-, -S- or -O-; and R², R³ and R⁴ are independently an optionally substituted aliphatic hydrocarbon group or an optionally substituted alicyclic heterocyclic ring group, or a pro-drug
- 10 thereof.
2. A compound according to claim 1, wherein R² and R³ are independently an optionally substituted acyclic hydrocarbon group.
3. A compound according to claim 1, wherein R² and R³ are
- 15 independently an optionally substituted alkyl group.
4. A compound according to claim 1, wherein R⁴ is an optionally substituted alicyclic hydrocarbon group or an optionally substituted alicyclic heterocyclic ring group.
5. A compound according to claim 4, wherein the alicyclic
- 20 hydrocarbon group is cycloalkyl.
6. A compound according to claim 4, wherein the alicyclic hydrocarbon group is cyclohexyl.
7. A compound according to claim 4, wherein the alicyclic heterocyclic ring group is a saturated alicyclic
- 25 heterocyclic ring group.
8. A compound according to claim 4, wherein the alicyclic heterocyclic ring group is tetrahydropyranyl, tetrahydrothiopyranyl or piperidyl.

9. N,N-Dimethyl-N-(4-(((2-(4-methylphenyl)-6,7-dihydro-5H-benzocyclohepten-8-yl)carbonyl)amino)benzyl)-N-(4-tetrahydropyranyl)ammonium chloride.
- 5 10. N,N-Dimethyl-N-(((7-(4-methylphenyl)-2,3-dihydro-1-benzoxepin-4-yl)carbonyl)amino)benzyl)-N-(4-oxocyclohexyl)ammonium chloride.
11. N-(4-(((7-(4-Ethoxyphenyl)-2,3-dihydro-1-benzoxepin-4-yl)carbonyl)amino)benzyl)-N,N-dimethyl-N-
- 10 (4-tetrahydropyranyl)ammonium chloride.
12. A pharmaceutical composition which comprises a compound as claimed in claim 1.
13. A pharmaceutical composition for antagonizing CCR5 which comprises a compound as claimed in claim 9.
- 15 14. A composition according to claim 13, which is for the treatment or prevention of infectious disease of HIV.
15. A composition according to claim 13, which is for the treatment or prevention of AIDS.
16. A composition according to claim 13, which is for the
- 20 prevention of the progression of AIDS.
17. A composition according to claim 13, which is used in combination with a protease inhibitor and/or a reverse transcriptase inhibitor.
18. A composition according to claim 17, wherein the
- 25 reverse transcriptase inhibitor is zidovudine, didanosine, zalcitabine, lamivudine, stavudine, nevirapine, delavirdine, efavirenz or abacavir.
19. A composition according to claim 17, wherein the protease inhibitor is saquinavir, ritonavir, indinavir or
- 30 nelfinavir.
20. Use of the compound as claimed in claim 9 in combination with a protease inhibitor and/or a reverse transcriptase inhibitor for the treatment or prevention of infectious disease of HIV.
- 35 21. Use of the compound as claimed in claim 9, for the manufacture of a medicament for antagonizing CCR5.

22. A method for antagonizing CCR5 which comprises administering to a mammal in need thereof an effective amount of the compound as claimed in claim 9.

SEQUENCE LIST

Sequence ID No. 1

Length of Sequence : 34

5 Type of Sequence : nucleic acid

Number of Chain : single

Topology : straight

Kind of Sequence : other nucleic acid synthetic DNA

Sequence:

10 CAGGATCCGA TGGATTATCA AGTGTCAAGT CCAA 34

Sequence ID No. 2

Length of Sequence : 34

Type of Sequence : nucleic acid

15 Number of Chain : single

Topology : straight

Kind of Sequence : other nucleic acid synthetic DNA

Sequence:

TCTAGATCAC AAGCCCACAG ATATTTCTTG CTCC 34

20